

BICYCLIC AROMATIC COMPOUNDS
AND PHARMACEUTICAL/COSMETIC
COMPOSITIONS COMPRISED THEREOF

CROSS-REFERENCE TO PRIORITY APPLICATION

5 The application claims priority under 35 U.S.C.
§ 119 of FR-98/03976, filed March 31, 1998, hereby
expressly incorporated by reference.

BACKGROUND OF THE INVENTION

Technical Field of the Invention:

10 The present invention relates to novel
biaromatic compounds and to pharmaceutical/cosmetic
compositions comprised thereof; the subject compounds
are especially useful in human or veterinary medicine,
or, alternatively, in cosmetic compositions.

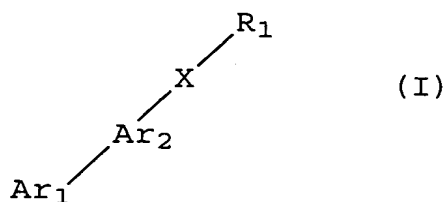
15 SUMMARY OF THE INVENTION

 The compounds according to the invention have
pronounced activity in the fields of cell
differentiation and proliferation, and are particularly
useful in the topical and systemic treatment of
20 dermatological conditions associated with a
keratinization disorder, dermatological (or other)

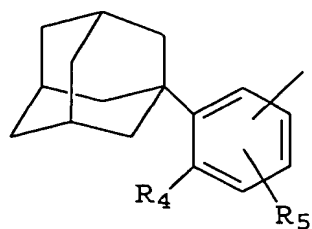
conditions including an inflammatory and/or immunoallergic component, and dermal or epidermal proliferations, whether benign or malignant. The subject compounds can also be used for the treatment of
5 degenerative diseases of connective tissue, to combat aging of the skin, whether photoinduced or chronological aging, and to treat cicatrization disorders. They are also useful for ophthalmological applications, especially for the treatment of corneopathies.

10 The compounds according to the invention can also be formulated into cosmetic compositions for body and hair hygiene.

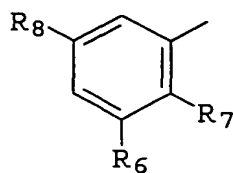
Briefly, the bicyclic aromatic compounds according to this invention have the following
15 structural formula (I):



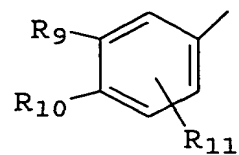
in which R_1 is (i) the radical $-\text{CH}_3$, (ii) a radical $-\text{CH}_2\text{OR}_2$, or (iii) a radical $-\text{CO}-\text{R}_3$, wherein R_2 and R_3 are as defined below; Ar_1 is a radical selected from among those of the following formulae (a)-(c):



(a)

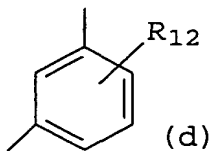


(b)

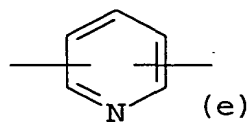


(c)

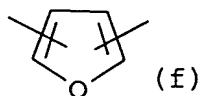
wherein R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} and R_{11} are as defined below; Ar_2 is a radical selected from among those of the following formulae (d) - (h):



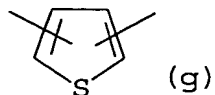
(d)



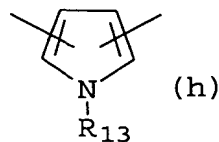
(e)



(f)

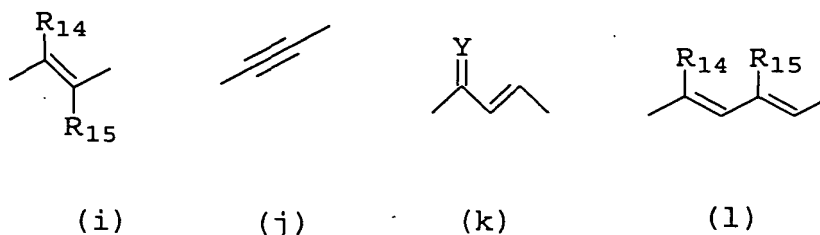


(g)

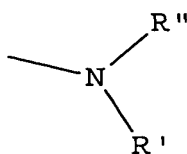


(h)

wherein R_{12} and R_{13} are as defined below; X is a radical
5 selected from among those of the following formulae
(i) - (l):



wherein R_{14} , R_{15} and Y are as defined below; R_2 is a hydrogen atom, a lower alkyl radical or a radical -CO- R_{16} , wherein R_{16} is as defined below; R_3 is a hydrogen atom, a lower alkyl radical, a radical of the
 5 formula:



wherein R' and R'' are as defined below, or a radical -OR₁₇, wherein R_{17} is as defined below; R_4 is a hydrogen atom, a polyether radical, a lower alkyl radical, or a radical -OR₁₈, wherein R_{18} is as defined below; R_5 is a
 10 hydrogen atom, a lower alkyl radical, a polyether radical, or a radical -OR₁₉, wherein R_{19} is as defined below; R_6 is a tert-butyl radical; R_7 is a lower alkyl radical, a polyether radical, or a radical -OR₂₀, wherein R_{20} is as defined below, with the proviso that R_6
 15 and R_7 may together form, with the carbon atoms from

which they depend, a 6-membered ring optionally substituted with at least one methyl group and/or optionally interrupted by an oxygen or sulfur atom; R_8 is a tert-butyl, adamantyl, aryl or aralkyl radical; R_9 and R_{10} together form, with the carbon atoms from which they depend, a 5- or 6-membered ring optionally substituted with at least one methyl group and/or optionally interrupted by an oxygen or sulfur atom; R_{11} is a hydrogen atom, a lower alkyl radical having from 1 to 9 carbon atoms, a hydroxyl radical, an alkoxy radical, a polyether radical or a radical $-OR_{21}$, wherein R_{21} is as defined below; R_{12} is a hydrogen atom, a hydroxyl radical, an alkoxy radical, a polyether radical or a radical $-OR_{22}$, wherein R_{22} is as defined below; R_{13} is a hydrogen atom, a lower alkyl radical, or a radical $-COR_{23}$, wherein R_{23} is as defined below; R_{14} and R_{15} , which may be identical or different, are each a hydrogen atom, or a lower alkyl radical; Y is an oxygen atom or a CH_2 radical; R_{16} is a lower alkyl radical; R_{17} is a hydrogen atom, a linear or branched alkyl radical having from 1 to 20 carbon atoms, an alkenyl radical, a mono- or polyhydroxyalkyl radical, an optionally substituted aryl or aralkyl radical or a sugar residue; R' and R'' , which may be identical or different, are each a hydrogen atom, a lower alkyl radical, a mono- or polyhydroxyalkyl radical, an optionally substituted aryl radical or an amino acid residue, with the proviso that R' and R'' may together form, with the nitrogen atom from which they

depend, a nitrogenous heterocycle; R_{18} , R_{19} and R_{20} , which may be identical or different, are each a hydrogen atom, a lower alkyl radical, an alkenyl radical, a mono- or polyhydroxyalkyl radical, an aryl radical, an optionally substituted aralkyl radical or a radical $-(CH_2)_n-R_{24}$, wherein n and R_{24} are as defined below; R_{21} and R_{22} , which may be identical or different, are each an alkenyl radical, a mono- or polyhydroxyalkyl radical, an aryl radical, an optionally substituted aralkyl radical, or a radical $-(CH_2)_n-R_{24}$, wherein n and R_{24} are as defined below; R_{23} is a lower alkyl radical; R_{24} is a heterocycle, a monohydroxyalkyl radical, a thiol radical optionally substituted with a lower alkyl radical, an amino radical optionally substituted with at least one lower alkyl radical, a radical $-COOR_{25}$, or a radical $-CON(R_{26})R_{27}$, wherein R_{25} , R_{26} and R_{27} are as defined below; R_{25} is a hydrogen atom or a lower alkyl radical; R_{26} and R_{27} , which may be identical or different, are each a hydrogen atom, a lower alkyl radical, or an optionally substituted aryl radical, with the proviso that R_{26} and R_{27} may together form, with the nitrogen atom from which they depend, a nitrogenous heterocycle; n is an integer such that $2 \leq n \leq 9$; with the further provisos that, when Ar_1 is the radical of formula (c) and X is a radical of formulae (i) or (j), then R_{11} is the radical $-OR_{21}$ or a polyether radical, or when Ar_2 is the radical of formula (d), then R_{12} is the radical $-OR_{22}$ or a polyether radical, and the compounds of formula (I)

do not include those wherein R_{11} is a methoxymethoxy radical in an ortho position relative to the substituent Ar_2 when Ar_1 is the radical of formula (c), X is the radical of formula (i), R_{12} is hydrogen and R_1 is the radical -COR₃ wherein R_3 is the radical -OR₁₇ and R_{17} is a hydrogen atom, and those wherein R_{12} is a methoxymethoxy radical in an ortho- or para-position relative to the substituent Ar_1 when Ar_1 is the radical of formula (c), X is the radical of formula (i), R_{11} is a methyl radical in an ortho-position relative to Ar_2 , and R_1 is the radical -COR₃ wherein R_3 is the radical -OR₁₇ and R_{17} is a hydrogen atom.

This invention also features the salts of the compounds of formula (I) when R_1 represents a carboxylic acid function, and the geometrical and optical isomers of said compounds of formula (I).

When the compounds according to the invention are in the form of salts, they are preferably salts of an alkali or alkaline earth metal, or, alternatively, of zinc or of an organic amine.

BRIEF DESCRIPTION OF THE DRAWINGS

The Figure of Drawing sets forth reaction schemes/mechanisms illustrating representative syntheses for the preparation of the bicyclic aromatic compounds according to the present invention.

DETAILED DESCRIPTION OF BEST MODE AND
SPECIFIC/PREFERRED EMBODIMENTS OF THE INVENTION

More particularly according to the present invention, by the term "lower alkyl radical" is intended
5 an alkyl radical having from 1 to 12, preferably from 1 to 9, carbon atoms, advantageously the methyl, ethyl, propyl, isopropyl, butyl, tert-butyl, pentyl, hexyl, heptyl, nonyl, decyl and dodecyl radicals.

By the expression "linear or branched alkyl
10 radical having from 1 to 20 carbon atoms" are particularly intended the methyl, ethyl, propyl, isopropyl, tert-butyl, octyl and 2-ethylhexyl radicals.

By the term "alkoxy radical" is intended an alkoxy radical preferably having from 1 to 9 carbon
15 atoms, in particular the methoxy, propyloxy, pentyloxy and heptyloxy radicals.

By the term "monohydroxyalkyl" or "polyhydroxyalkyl" radical is intended a radical having from 1 to 6 carbon atoms and from 1 to 5 hydroxyl
20 groups. Among such monohydroxyalkyl radicals, preferred are those having 2 or 3 carbon atoms, in particular a 2-hydroxyethyl, 2-hydroxypropyl or 3-hydroxypropyl radical. Among such polyhydroxyalkyl radicals, preferred are those having from 3 to 6 carbon atoms and
25 from 2 to 5 hydroxyl groups, such as the 2,3-dihydroxypropyl, 2,3,4-trihydroxybutyl and 2,3,4,5-tetrahydroxypentyl radicals, or the pentaerythritol residue.

By the term "aryl radical" is preferably intended a phenyl radical optionally substituted with at least one halogen atom, lower alkyl radical, hydroxyl group, alkoxy radical, nitro function, polyether radical, or an amino function optionally protected with an acetyl group or optionally substituted with at least one lower alkyl radical.

By the term "aralkyl radical" is preferably intended a benzyl or phenethyl radical optionally substituted with at least one halogen atom, lower alkyl radical, hydroxyl group, alkoxy radical, nitro function, a polyether radical or an amino function optionally protected with an acetyl group or optionally substituted with at least one lower alkyl radical.

By the term "alkenyl radical" is preferably intended an alkenyl radical having from 2 to 5 carbon atoms and containing one or more sites of ethylenic unsaturation, such as, more particularly, the allyl radical.

By the term "polyether radical" is intended a radical having from 1 to 6 carbon atoms and from 1 to 3 oxygen or sulfur atoms, such as the methoxymethyl ether, methoxyethoxymethyl ether or methylthiomethyl ether radicals.

By "sugar residues" are preferably intended those derived, for example, from glucose, galactose, mannose or glucuronic acid.

By "amino acid residue" is preferably intended one derived, for example, from at least one of the 20 amino acids of L or D configuration constituting mammalian proteins. More particularly intended are the residues derived from lysine, glycine or aspartic acid.

By "heterocycle" is preferably intended piperidino, morpholino, pyrrolidino or piperazino, optionally substituted in the 4-position with a C₁-C₆ alkyl radical or with a mono- or polyhydroxyalkyl radical as defined above.

Among the compounds of formula (I) according to the present invention, particularly representative are the following:

- 3-(3',5'-Di-tert-butyl-2'-methoxy-3-biphenyl)acrylic acid;
- 3-(2'-Benzyloxy-3',5'-di-tert-butyl-6-hydroxy-3-biphenyl)acrylic acid;
- 3-(3',5'-Di-tert-butyl-6-hydroxy-2'-pentyloxy-3-biphenyl)acrylic acid;
- 3-(3',5'-Di-tert-butyl-6-hydroxy-2'-methoxy-3-biphenyl)acrylic acid;
- 3-[3-(3-Benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid;
- 3-[3-(3-Benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-hydroxyphenyl]acrylic acid;
- [4-Methoxymethoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]propynoic acid;

3-[3-(3-Propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid;

5 3-[3-(3-Pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid;

3-(5'-Adamantan-1-yl-4'-methoxy-2'-methyl-3-biphenyl)acrylic acid;

10 3-(5'-Adamantan-1-yl-6-hydroxy-4'-methoxy-2'-methyl-3-biphenyl)acrylic acid;

3-(5'-Adamantan-1-yl-4'-methoxy-6-methoxymethoxy-2'-methyl-3-biphenyl)acrylic acid;

15 3-{4-Methoxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid;

3-{4-Methoxy-3-[3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid;

20 3-{3-[3-(6-Hydroxyhexyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

3-{3-[3-(7-Hydroxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

25 3-{3-[3-(5-Hydroxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

3-{3-[3-(3-Hydroxypropyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

5 3-[3-(1-Benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid;

3-(3'-Adamantan-1-yl-4'-hydroxy-3-biphenyl)acrylic acid;

10 5-[4-Methoxy-3-(5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]-3-methylpenta-2,4-dienoic acid;

5-[4-Methoxymethoxy-3-(5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]-3-methylpenta-2,4-dienoic acid;

15 5-[4-Hydroxy-3-(5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]-3-methylpenta-2,4-dienoic acid;

3-{3-[3-(5-Tert-butoxycarbonylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

20 3-{3-[3-(7-Tert-butoxycarbonylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

25 3-{3-[3-(7-Carboxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

3-{3-[3-(5-Carboxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

3-{3-[3-(5-Carbamoylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

5 3-{3-[3-(7-Carbamoylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid;

3-{4-Methoxy-3-[5,5,8,8-tetramethyl-3-(2-morpholin-4-ylethoxy)-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid;

10 3-{4-Methoxy-3-[5,5,8,8-tetramethyl-3-(2-piperidin-1-ylethoxy)-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid;

15 3-{4-Methoxy-3-[3-(2-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid;

3-{4-Methoxy-3-[3-(3-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid;

20 3-{4-Methoxy-3-[3-(4-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid;

3-{4-Methoxy-3-[3-(3-hydroxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid;

25 3-[4-Fluoro-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid;

3-{4-Hydroxy-3-[3-(3-methoxybenzyloxy)-
5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-
naphthyl]phenyl}acrylic acid;

5 3-{4-Hydroxy-3-[3-(4-methoxybenzyloxy)-
5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-
naphthyl]phenyl}acrylic acid;

3-{4-Hydroxy-3-[3-(4-fluorobenzyloxy)-5,5,8,8-
tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic
acid;

10 3-{4-Hydroxy-3-[3-(4-chlorobenzyloxy)-5,5,8,8-
tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic
acid;

15 3-[4-Hydroxy-3-(3-methoxymethoxy-5,5,8,8-
tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic
acid;

3-(3',5'-Di-tert-butyl-6-hydroxy-2'-propyloxy-
3-biphenyl)acrylic acid;

3-(3',5'-Di-tert-butyl-6-hydroxy-2'-butyloxy-
3-biphenyl)acrylic acid;

20 3-(2'-Butoxy-3',5'-di-tert-butyl-6-methoxy-3-
biphenyl)acrylic acid;

3-(3'5'-Di-tert-butyl-6-methoxy-2'-propoxy-3-
biphenyl)acrylic acid;

25 3-[4-Hydroxy-3-(5,5,8,8-tetramethyl-4-propoxy-
5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid;

3-[3-(3-Benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-
tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylic acid;

Ethyl 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate;

3-Methyl-5-[3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]penta-2,4-dienoic acid;

5 3-[3-(3-Benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]prop-2-en-1-ol;

3-[3-(3-Benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]propenal;

N-Ethyl 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]-acrylamide;

3-[3-(3-Benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]-1-morpholin-4-yl-propenone;

15 N-(4-Hydroxyphenyl)-3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylamide;

5-(5'-Adamantan-1-yl-4'-methoxy-2'-methyl-3-biphenyl)-3-methylpenta-2,4-dienoic acid;

20 5-[4-Methoxymethoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]-3-methylpenta-2,4-dienoic acid;

5-[4-Hydroxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]-3-methylpenta-2,4-dienoic acid;

25 4-[3-(3,5,5,8,8-Pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]penta-2,4-dienoic acid;

5-(3',5'-Di-tert-butyl-2'-methoxy-3-biphenylyl)-3-methylpenta-2,4-dienoic acid;

5-(3',5'-Di-tert-butyl-2'-propoxy-3-biphenylyl)-3-methylpenta-2,4-dienoic acid;

5 5-(2'-Butoxy-3',5'-di-tert-butyl-3-biphenylyl)-3-methylpenta-2,4-dienoic acid;

5-(2'-Butoxy-3',5'-di-tert-butyl-6-hydroxy-3-biphenylyl)-3-methylpenta-2,4-dienoic acid;

10 5-(3',5'-Di-tert-butyl-6-hydroxy-2'-propoxy-3-biphenylyl)-3-methylpenta-2,4-dienoic acid;

5-(3',5'-Di-tert-butyl-6-hydroxy-2'-methoxy-3-biphenylyl)-3-methylpenta-2,4-dienoic acid.

According to the present invention, the compounds of formula (I) which are more particularly
15 preferred are those for which at least one, and preferably all, of the following conditions are satisfied:

- 20 R_1 is a radical $-CO-R_3$;
 Ar_2 is a radical of formulae (d) or (e);
 R_{11} is a radical $-O-R_{21}$;
 R_7 is a radical $-O-R_{20}$.

The present invention also features processes for preparing the compounds of formula (I), in particular according to the reaction scheme illustrated
25 in the Figure of Drawing.

The compounds of formula (Ia) can be obtained (Figure of Drawing) from aldehyde or ketone derivatives (5), according to a Horner-type reaction with a lithium

or sodium derivative of a phosphonate (7), it being possible for the carbonyl compounds (5) to be obtained either by a coupling reaction between a boronic acid (3) and a halo derivative (4) (this reaction is carried out in the presence of a palladium catalyst, for example tetrakis(triphenylphosphine)-palladium under the conditions described by N. Miyaura et al., Synthetic Communications, 11(7), 513-519 (1981); the boronic acid derivative (3) can be obtained, for example, from the halo derivative (1) by conversion into the lithium derivative (2) and then reaction with trimethyl borate and hydrolysis), or by a coupling reaction between a zinc derivative (8) and a halo ester derivative (9) in the presence of a catalyst, for example a palladium or nickel derivative (NiCl_2dppe), followed by conversion of the ester function (10) into an alcohol (11) and oxidation into an aldehyde (5).

The compounds of formula (Ib) can be obtained from the acetylenic derivative (6) by reaction with n-butyllithium and then carboxylation in the presence of CO_2 .

The acetylenic compounds (6) can be obtained either:

from the aldehyde derivatives (5) (when R_3 is a hydrogen atom), by reaction with carbon tetrabromide and triphenylphosphine to give a 2'2'-dibromostyrene derivative which is converted into an acetylenic derivative by a non-nucleophilic base such as n-

butyllithium in an aprotic solvent such as tetrahydrofuran, or

from the ketone derivatives (5) (when R_3 is a lower alkyl radical) by a sequence of reactions comprising treatment with a base such as lithium diisopropylamide, then with a dialkyl phosphate chloride and again with lithium diisopropylamide.

When R_1 is a $-COOH$ radical, the compounds are prepared:

either by protecting R_1 with a protecting group of alkyl, allylic or tert-butyl type.

Conversion to the free form can be carried out:

in the case of an alkyl protecting group, using sodium hydroxide or lithium hydroxide in an alcoholic solvent such as methanol or in THF;

in the case of an allylic protecting group, using a catalyst such as certain transition metal complexes in the presence of a secondary amine such as morpholine;

in the case of a protecting group of tert-butyl type, using trimethylsilane iodide.

When R_1 is an alcohol function, the compounds can be obtained:

from the corresponding aldehyde derivatives by the action of an alkyl metal hydride, such as sodium borohydride, in an alcoholic solvent (for example methanol);

from the acid derivatives by reduction with lithium aluminum hydride.

When R_1 is an aldehyde function, the compounds can be obtained from the alcohol derivatives by
5 oxidation in the presence of manganese oxide, pyridinium dichromate or Swern's reagent.

When R_1 is an amide function, the compounds can be obtained from the corresponding carboxylic derivatives by reaction with aliphatic, aromatic or
10 heterocyclic amines either via an acid chloride or in the presence of dicyclohexylcarbodiimide or carbonyldiimidazole.

Certain of the subject compounds bind to the RXR receptors, some having agonist activity, others
15 having antagonist activity. The preferred compounds have antagonist activity.

The binding and transactivation properties as RXR-receptor agonists can be determined by methods known in the art, such as, for example: Levin et al., Nature,
20 355, 359-61 (1992); Allenby et al., Proc. Natl. Acad. Sci., 90, 30-4 (1993).

The RXR-agonist activity can also be determined by the test described in French patent application No. 95/07301, filed June 19, 1995 by the
25 assignee hereof. This test comprises the following steps: (i) a sufficient amount of a compound which is an active ligand for at least one receptor of the superfamily of steroidal/thyroid nuclear receptors other

than a ligand specific for the RXR receptors and which can heterodimerize with the RXRs, such as an RAR-agonist molecule, is applied topically to an area of the skin of a mammal, (ii) a molecule capable of having RXR-agonist activity is administered systemically or topically to this same area of the skin of the mammal before, during or after step (i), and (iii) the response on the part of the mammal's skin thus treated is evaluated. Thus, the response to topical application to a mammal's ear of an RAR-agonist molecule which corresponds to an increase in the thickness of this ear can be increased by the systemic or topical administration of an RXR-receptor agonist molecule.

The RXR α -antagonist activity can be evaluated in the transactivation test by determination of the dose (IC₅₀) which gives 50% inhibition of the transactivating activity of an RXR α -selective agonist: 6-[1-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)cyclopropyl]nicotinic acid (CD 3127) according to the following procedure:

HeLa cells are co-transfected with an expression vector coding for RXR α (p565-RXR α) and a reporter plasmid containing the response element $\frac{1}{2}$ CRBP II cloned upstream of the thymidine kinase heterologous promoter and of the chloramphenicolm-acetyl-transferase (CAT) reporter gene. Eighteen hours after co-transfection, the cells are treated with a fixed concentration of CD 3127 and increasing concentrations

of the molecule to be evaluated. After treatment for twenty-four hours, the CAT activity is assayed by ELISA. The fixed concentration of CD3127 used is 10^{-8} M and corresponds to its EC₅₀.

5 The compounds according to the invention are particularly well suited in the following fields of therapy:

 (1) for treating dermatological conditions associated with a keratinization disorder related to
10 differentiation and on proliferation, in particular for treating common acne, comedones, polymorphonuclear leukocytes, rosacea, nodulocystic acne, acne conglobata, senile acne and secondary acnes such as solar, medication-related or occupational acne;

15 (2) for treating other types of keratinization disorders, in particular ichthyosis, ichthyosiform states, Darier's disease, palmoplantar keratoderma, leucoplasias and leucoplasiform states, and cutaneous or mucous (buccal) lichen;

20 (3) for treating other dermatological conditions associated with a keratinization disorder manifesting an inflammatory and/or immunoallergic component and, in particular, all forms of psoriasis, whether cutaneous, mucous or unguual psoriasis and even
25 psoriatic rheumatism, or, alternatively, cutaneous atopy, such as eczema or respiratory atopy, or gingival hypertrophy; the compounds can also be used for treating

inflammatory conditions not exhibiting keratinization disorder;

(4) for treating all dermal or epidermal proliferations, whether benign or malignant and whether
5 or not of viral origin or otherwise, such as common warts, flat warts and verruciform epidermodysplasia, oral or florid papillomatoses and the proliferations which can be induced by ultraviolet radiation, in particular in the case of basocellular and spinocellular
10 epithelioma;

(5) for treating other dermatological disorders such as bullosis and collagen diseases;

(6) for treating certain ophthalmological disorders, in particular corneopathies;

15 (7) for repairing or combating aging of the skin, whether photoinduced or chronological, or for reducing actinic keratoses and pigmentations, or any pathologies associated with chronological or actinic aging;

20 (8) for preventing or curing the stigmata of epidermal and/or dermal atrophy induced by local or systemic corticosteroids, or any other form of cutaneous atrophy;

(9) for preventing or treating cicatrization
25 disorders or for preventing or repairing stretch marks, or, alternatively, for promoting cicatrization;

(10) for combating disorders of sebaceous functioning, such as the hyperseborrhoea of acne or simple seborrhoea;

5 (11) for the treatment or prevention of cancerous or precancerous states;

(12) for the treatment of inflammatory conditions such as arthritis;

(13) for the treatment of any general or skin affliction of viral origin;

10 (14) for the prevention or treatment of alopecia;

(15) for the treatment of dermatological or general conditions including an immunological component;

15 (16) for the treatment of conditions of the cardiovascular system such as arteriosclerosis, hypertension, non-insulin-dependent diabetes and obesity;

(17) for the treatment of skin disorders due to an exposure to U.V. radiation.

20 In the therapeutic fields indicated above, the compounds according to the invention may advantageously be administered in combination with other compounds exhibiting retinoid-type activity, with D vitamins or derivatives thereof, with corticosteroids, with anti-
25 free-radical agents, with α -hydroxy or α -keto acids or derivatives thereof, or, alternatively, with ion-channel blockers. By the expression "D vitamins or derivatives thereof" are intended, for example, vitamin D₂ or D₃

derivatives and in particular 1,25-dihydroxyvitamin D₃.
By "anti-free-radical agents" are intended, for example,
 α -tocopherol, superoxide dismutase, ubiquinol or certain
metal-chelating agents. By the expression " α -hydroxy or
5 α -keto acids or derivatives thereof" are intended, for
example, lactic, malic, citric, glycolic, mandelic,
tartaric, glyceric or ascorbic acid or the salts, amides
or esters thereof. Lastly, by the term "ion-channel
blockers" are intended, for example, Minoxidil (2,4-
10 diamino-6-piperidinopyrimidine 3-oxide) and derivatives
thereof.

The present invention thus also features
medicinal compositions containing at least one compound
of formula (I), one of the optical or geometrical
15 isomers thereof, or one of the salts or other
derivatives thereof.

The pharmaceutical/therapeutic compositions of
this invention, intended especially for the treatment of
the aforesaid disease states comprise a carrier, diluent
20 or vehicle which is pharmaceutically acceptable and
compatible with the mode of administration selected for
the given composition, at least one compound of formula
(I), one of the optical or geometrical isomers thereof,
or one of the salts, etc., thereof.

25 The compounds according to the invention can
be administered systemically, enterally, parenterally,
topically or ocularly.

For enteral administration, the medicinal/pharmaceutical compositions may be in the form of tablets, gelatin capsules, sugar-coated tablets, syrups, suspensions, solutions, elixirs, powders, granules, emulsions, microspheres or nanospheres or polymeric or lipid vesicles which permit a controlled release. For parenteral administration, the compositions may be in the form of solutions or suspensions for perfusion or for injection.

10 The compounds according to the invention are generally administered at a daily dose of about 0.01 mg/kg to 100 mg/kg of body weight, and this at the regime or rate of 1 to 3 doses per diem.

 For topical administration, the pharmaceutical compositions based on compounds according to the invention are more particularly intended for the treatment of the skin and the mucosae and may be in the form of ointments, creams, milks, salves, powders, pommades, impregnated pads, solutions, gels, sprays, lotions or suspensions. They may also be in the form of microspheres or nanospheres or polymeric or lipid vesicles or polymeric patches and hydrogels which permit controlled release. These topical-route compositions may either be in anhydrous form or in aqueous form, depending on the clinical indication.

25 For ocular administration, they are mainly eyedrops.

These compositions for topical or ocular application contain at least one compound of formula (I) as defined above, or one of the optical or geometrical isomers thereof, or one of the salts or other
5 derivatives thereof, at a concentration preferably ranging from 0.001% to 5% by weight relative to the total weight of the composition.

The compounds of formula (I) according to the invention also find application in the cosmetic field,
10 in particular for body and hair hygiene and especially for treating skin types with a tendency towards acne, for promoting the regrowth of the hair, for combating hair loss, for combating the greasy appearance of the skin or the hair, for protection against the harmful
15 effects of the sun or for the treatment of physiologically dry skin types, and for preventing and/or combating photoinduced or chronological aging.

For cosmetic applications, the compounds according to the invention can moreover be employed
20 advantageously in combination with other compounds of retinoid-type activity, with D vitamins or derivatives thereof, with corticosteroids, with anti-free-radical agents, with α -hydroxy or α -keto acids or derivatives thereof, or alternatively with ion-channel blockers, all
25 of these various active agents being as defined above.

The present invention therefore also features cosmetic compositions comprising a cosmetically acceptable support (vehicle, diluent or carrier) which

is suitable for topical application, at least one compound of formula (I) as defined above or one of the optical or geometrical isomers thereof, or one of the salts or other derivatives thereof. Such cosmetic
5 compositions are advantageously in the form of a cream, a milk, a lotion, a gel, microspheres or nanospheres or polymeric or lipid vesicles, a soap or a shampoo.

The concentration of the compounds of formula (I) in the cosmetic compositions according to the
10 invention advantageously ranged from 0.001% to 3% by weight relative to the total weight of the composition.

The medicinal and cosmetic compositions of the present invention can also contain inert additives and adjuvants, or even pharmacodynamically or cosmetically
15 active additives and adjuvants, or combinations of these additives and adjuvants, in particular: wetting agents; depigmenting agents such as hydroquinone, azelaic acid, caffeic acid or kojic acid; emollients; moisturizing agents such as glycerol, PEG 400, thiamorpholinone and
20 derivatives thereof, or urea; anti-seborrhoea or anti-acne agents such as S-carboxymethylcysteine, S-benzylcysteamine, the salts and the derivatives thereof, or benzoyl peroxide; antibiotics such as erythromycin and esters thereof, neomycin, clindamycin
25 and esters thereof, and tetracyclines; antifungal agents such as ketoconazole or 4,5-polymethylene-3-isothiazolinones; agents for promoting the regrowth of the hair, such as Minoxidil (2,4-diamino-6-

piperidinopyrimidine 3-oxide) and derivatives thereof, Diazoxide (7-chloro-3-methyl-1,2,4-benzothiadiazine 1,1-dioxide) and Phenytoin (5,5-diphenylimidazolidine-2,4-dione); non-steroidal anti-inflammatory agents; carotenoids and, in particular, β -carotene; anti-psoriatic agents such as anthraline and derivatives thereof and, lastly, eicosa-5,8,11,14-tetraynoic acid and eicosa-5,8,11-triynoic acid, the esters and the amides thereof.

10 The compositions according to the invention may also contain flavor-enhancing agents, preservatives such as para-hydroxybenzoic acid esters, stabilizing agents, moisture regulators, pH regulators, osmotic pressure modifiers, emulsifying agents, UV-A and UV-B
15 screening agents, and antioxidants such as α -tocopherol, butylhydroxyanisole or butylhydroxytoluene.

 In order to further illustrate the present invention and the advantages thereof, the following specific examples are given, it being understood that
20 same are intended only as illustrative and in nowise limitative.

A. EXAMPLES OF SYNTHESSES OF COMPOUNDS, PER SE:

EXAMPLE 1:

Synthesis of 3-(3',5'-di-tert-butyl-2'-methoxy-3-biphenyl)acrylic acid:
25

(a) Preparation of 2-bromo-4,6-di-tert-butylanisole:

6 g (21 mmol) of 2-bromo-4,6-di-tert-butylphenol and 60 ml of DMF were introduced into a round-bottomed flask under a stream of nitrogen. 694 mg (29 mmol) of sodium hydride (80% in oil) were added portionwise and the mixture was stirred until the evolution of gas had ceased. 1.45 ml (23.2 mmol) of iodomethane was then added and the mixture was stirred at room temperature for three hours. The reaction medium was poured into ice-cold water and extracted with ethyl acetate, and the organic phase was separated out after settling had taken place, dried over magnesium sulfate and evaporated. The residue obtained was purified by chromatography from a column of silica eluted with heptane. After evaporation of the solvents, 6.1 g (98%) of the expected compound were collected in the form of a colorless oil.

^1H NMR (d, CDCl_3): 1.3 (s, 9H); 1.4 (s, 9H); 3.9 (s, 3H); 7.3 (d, $J = 2.3$ Hz, 1H); 7.4 (d, $J = 2.3$ Hz, 1H).

(b) Preparation of 2-methoxy-3,5-di-tert-butylphenylboronic acid:

6 g (20.1 mmol) of 2-bromo-4,6-di-tert-butylanisole and 50 ml of THF were introduced into a three-necked flask under a stream of nitrogen. 9.7 ml (24.1 mmol) of n-butyllithium (2.5M in hexane) were

added dropwise, at -78°C , and the mixture was stirred for 15 minutes. At this same temperature, 7 ml (30.1 mmol) of triisopropyl borate were added with stirring for 2 hours. 20 ml of hydrochloric acid (1N) were added at -50°C and the mixture was permitted to warm to room temperature. The reaction medium was extracted with ethyl ether and the organic phase was separated out after settling had taken place, dried over magnesium sulfate and evaporated. The solid obtained was triturated from heptane, filtered off and dried. 4 g (75%) of the expected boronic acid were collected, which product was used without further purification in the remainder of the synthesis to follow.

^1H NMR (d, CDCl_3): 1.3 (s, 9H); 1.4 (s, 9H); 3.8 (s, 3H); 6.0 (s, 2H); 7.5 (d, $J = 2.5$ Hz, 1H); 7.7 (d, $J = 2.5$ Hz, 1H).

(c) Synthesis of 3-(3',5'-di-tert-butyl-2'-methoxy-3-biphenyl)acrylic acid:

2.2 g (8.3 mmol) of 2-methoxy-3,5-di-tert-butylphenylboronic acid, 942 mg (4.15 mmol) of 3-bromocinnamic acid and 70 ml of DME were introduced into a three-necked flask under a stream of nitrogen. 11.4 ml of aqueous potassium carbonate solution (2M) were added dropwise and the reaction medium was degassed. 144 mg (0.12 mmol) of tetrakis(triphenylphosphine)palladium(0) were then added and the mixture was refluxed for 20

hours. Water and ethyl acetate were added to the reaction medium and this mixture was acidified to pH 1 with hydrochloric acid. The organic phase was separated out after settling had taken place, washed with water, 5 dried over magnesium sulfate and evaporated. The residue obtained was purified by chromatography on a column of silica eluted with a mixture of heptane and ethyl acetate (75/25). After evaporation of the solvents, 990 mg (66%) of 3-(3',5'-di-tert-butyl-2'-methoxy-3- 10 biphenylyl)acrylic acid having a melting point of 182°-3°C were collected.

¹H NMR (d, DMSO): 1.1 (s, 9H); 1.2 (s, 9H); 3.0 (s, 3H); 6.4 (d, J = 16 Hz, 1H); 7.0 (d, J = 2.3 Hz, 1H); 7.1 (d, J = 2.3 Hz, 1H); 7.3-7.6 (m, 5H).

15

EXAMPLE 2:

Synthesis of 3-(2'-benzyloxy-di-tert-butyl-6-hydroxy-3-biphenylyl)acrylic acid:

(a) Preparation of 3-bromo-4-hydroxybenzaldehyde:

20 15 g (123 mmol) of 4-hydroxybenzaldehyde, 687 mg (12.3 mmol) of iron powder, 200 ml of dichloromethane and 10 ml of THF were introduced into a 500 ml three-necked flask under nitrogen atmosphere. The mixture was cooled to 0°C, 19.60 g (123 mmol) of bromine 25 were added dropwise and this mixture was stirred at room

temperature for one hour. Saturated sodium thiosulfate solution was added, the dichloromethane was evaporated off, the residue was extracted with ethyl acetate, the organic phase was separated out after settling had taken place, washed with water to neutral pH and dried over magnesium sulfate, and the solvents were evaporated off. The residue obtained was purified by chromatography on a column of silica eluted with an eluting system composed of 80% heptane and 20% dichloromethane. After evaporation of the solvents, 9.7 g (39%) of a white cottony powder having a melting point of 121°-124°C were collected.

(b) Preparation of 3-bromo-4-methoxymethoxybenzaldehyde:

9 g (44.7 mmol) of 3-bromo-4-hydroxybenzaldehyde, 30 ml of THF and 30 ml of DMF were introduced into a 250 ml round-bottomed flask under a nitrogen atmosphere. 1.48 g (49.2 mmol) of 80% sodium hydride were added portionwise, the mixture was stirred for thirty minutes at room temperature and 4.32 g (52.7 mmol) of methoxymethyl chloride were added dropwise. The reaction medium was stirred for one hour at room temperature, acidified with 1N hydrochloric acid and extracted with ethyl acetate. The organic phase was separated out after settling had taken place, washed with water to neutral pH and dried over magnesium

sulfate, and the solvents were evaporated off. 11.5 g (100%) of an orange-colored oil were collected.

(c) Preparation of ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate:

5 12 g (53 mmol) of triethyl phosphonoacetate
and 50 ml of THF were introduced into a three-necked
flask under a stream of nitrogen. 2.45 g (81.6 mmol) of
sodium hydride (80% in oil) were added portionwise and
the mixture was stirred until the evolution of gas had
10 ceased. A solution of 10 g (40.8 mmol) of 3-bromo-4-
methoxymethoxybenzaldehyde in 50 ml of THF was then
added and this mixture was stirred at room temperature
for one hour. The reaction medium was poured into water,
acidified with hydrochloric acid and extracted with
15 ethyl acetate, and the organic phase was separated out
after settling had taken place, dried over magnesium
sulfate and evaporated. The residue obtained was
purified by chromatography on a column of silica eluted
with a mixture of heptane and ethyl acetate (90-10).
20 After evaporation of the solvents, 9.6 g (74%) of ethyl
3-(3-bromo-4-methoxymethoxyphenyl)acrylate having a
melting point of 63°-65°C were collected.

(d) Preparation of 2-benzyloxy-3,5-di-tert-butylbromobenzene:

In a manner similar to that of Example 1(a),
by reaction of 6 g (21 mmol) of 2-bromo-4,6-di-tert-
5 butylphenol with 2.8 ml (23.2 mmol) of benzyl bromide,
7.4 g (94%) of the expected compound were obtained in
the form of a colorless oil.

¹H NMR (d, CDCl₃): 1.3 (s, 9H); 1.4 (s, 9H); 5.1 (s,
2H); 7.25-7.45 (m, 5H); 7.55-7.6 (m, 2H).

10 (e) Preparation of 2-benzyloxy-3,5-di-tert-butylphenylboronic acid:

In a manner similar to that of Example 1(b),
by reaction of 7.2 g (19.3 mmol) of 2-benzyloxy-3,5-di-
tert-butylbromobenzene with 6.7 ml (29 mmol) of
15 triisopropyl borate, 4.29 g (65%) of the expected
boronic acid were obtained.

¹H NMR (d, CDCl₃): 1.35 (s, 9H); 1.45 (s, 9H); 5.1 (s,
2H); 5.9 (s, 2H); 7.3-7.55 (m, 6H); 7.7 (d, J = 2.5 Hz,
1H).

20 (f) Preparation of ethyl 3-(2'-benzyloxy-3',5'-di-tert-butyl-6'-methoxymethoxy-3-biphenyl)acrylate:

In a manner similar to that of Example 1(c),
by reaction of 3.24 g (9.53 mmol) of 2-benzyloxy-3,5-di-

tert-butylphenylboronic acid with 2 g (6.35 mmol) of ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate, 2.4 g (71%) of the expected ethyl ester were obtained.

¹H NMR (d, CDCl₃): 1.1 (t, J = 7 Hz, 3H); 1.35 (s, 9H);
5 1.45 (s, 9H); 3.5 (s, 3H); 4.25 (q, J = 7 Hz, 2H); 4.5 (s, 2H); 5.1 (s, 2H); 6.3 (d, J = 16 Hz, 1H); 6.9-7.0 (m, 2H); 7.15-7.25 (m, 5H); 7.4-7.5 (m, 2H); 7.6-7.7 (m, 2H).

10 (g) Preparation of ethyl 3-(2'-benzyloxy-3',5'-di-tert-butyl-6-hydroxy-3-biphenyl)acrylate:

2.14 g (4 mmol) of the above ethyl ester, 10 ml of THF and 10 ml of ethanol were introduced into a round-bottomed flask. 1 ml of concentrated sulfuric acid was added and the mixture was stirred at room
15 temperature for 24 hours. The reaction medium was poured into water and extracted with ethyl acetate, and the organic phase was separated out after settling had taken place, washed with water, dried over magnesium sulfate and evaporated. The residue obtained was triturated from
20 heptane, filtered and dried. 1.93 g (100%) of the expected compound was collected.

¹H NMR (d, CDCl₃): 1.3 (t, J = 7 Hz, 3H); 1.35 (s, 9H); 1.45 (s, 9H); 4.25 (q, J = 7 Hz, 2H); 4.5-4.7 (m, 2H); 6.4 (d, J = 16 Hz, 1H); 7.0 (d, J = 8 Hz, 1H); 7.0-7.1
25 (m, 2H); 7.25 (s, 1H); 7.3-7.4 (m, 3H); 7.45-7.50 (m, 2H); 7.6 (s, 1H); 7.7 (d, J = 6 Hz, 1H).

(h) Synthesis of 3-(2'-benzyloxy-3',5'-di-tert-butyl-6-hydroxy-3-biphenyl)acrylic acid:

1.93 g (4 mmol) of ethyl 3-(2'-benzyloxy-3',5'-di-tert-butyl-6-hydroxy-3-biphenyl)acrylate,
5 20 ml of THF, 2 ml of methanol and 4 ml of sodium hydroxide solution (10N) were introduced into a round-bottomed flask. The reaction medium was refluxed for 12 hours, poured into water, acidified to pH 1 with hydrochloric acid and extracted with ethyl acetate, and
10 the organic phase was separated out after settling had taken place, dried over magnesium sulfate and evaporated. The residue obtained was triturated from heptane, filtered and dried. 1.6 g (87%) of 3-(2'-benzyloxy-3',5'-di-tert-butyl-6-hydroxy-3-biphenyl)acrylic acid having a melting point of 230°-1°C was collected.
15 ¹H NMR (d, CDCl₃): 1.3 (s, 9H); 1.5 (s, 9H); 4.6 (s, 2H); 6.3 (d, J = 16 Hz, 1H); 7.0-7.1 (m, 3H); 7.2-7.3 (m, 4H); 7.4-7.5 (m, 2H); 7.6 (s, 1H); 7.7 (d, J = 16
20 Hz, 1H); 8.1 (m, 1H).

EXAMPLE 3:

Synthesis of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-pentyloxy-3-biphenyl)acrylic acid:

(a) Synthesis of 3,5-di-tert-butyl-2-pentyloxybromobenzene:

In a manner similar to that of Example 1(a),
by reaction of 6 g (21 mmol) of 2-bromo-4,6-di-tert-
5 butylphenol with 3 ml (23.2 mmol) of iodopentane, 7.3 g
(98%) of the expected compound were obtained in the form
of a colorless oil.

¹H NMR (d, CDCl₃): 0.9 (t, J = 7 Hz, 3H); 1.35 (s, 9H);
1.45 (s, 9H); 1.3-1.5 (m, 4H); 1.8-2.0 (m, 2H); 4.0 (q,
10 J = 7 Hz, 2H); 7.3 (d, J = 2.5 Hz, 1H); 7.4 (d, J = 2.5
Hz, 1H).

(b) Preparation of 3,5-di-tert-butyl-2-pentyloxyphenylboronic acid:

In a manner similar to that of Example 1(b),
15 by reaction of 7.26 g (20.5 mmol) of 3,5-di-tert-butyl-
2-pentyloxybromobenzene with 7.1 ml (30.75 mmol) of
triisopropyl borate, 6.4 g (98%) of the expected boronic
acid were obtained.

¹H NMR (d, CDCl₃): 0.9 (t, J = 7 Hz, 3H); 1.2-1.5 (m,
20 4H); 1.35 (s, 9H); 1.45 (s, 9H); 1.85-1.95 (m, 2H); 3.8-
4.0 (m, 2H); 6.1 (s, 1H); 6.8-7.6 (m, 3H).

(c) Preparation of ethyl 3-(3',5'-di-tert-butyl-6-methoxymethoxy-2'-pentyloxy-3-biphenyl)acrylate:

In a manner similar to that of Example 1(c),
5 by reaction of 3 g (9.5 mmol) of 3,5-di-tert-butyl-2-pentyloxyphenylboronic acid with 2 g (6.35 mmol) of ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate, 2.1 g (65%) of the expected ethyl ester are obtained.

¹H NMR (d, CDCl₃): 0.7 (t, J = 7 Hz, 3H); 1.0 (m, 4H);
10 1.2 (m, 5H); 1.25 (s, 9H); 1.35 (s, 9H); 3.2-3.3 (m, 2H); 3.35 (s, 3H); 4.2 (q, J = 7 Hz, 2H); 5.1 (s, 2H); 6.25 (d, J = 16 Hz, 1H); 7.05 (d, J = 2.5 Hz, 1H); 7.15 (d, J = 8.5 Hz, 1H); 7.3 (d, J = 2.5 Hz, 1H); 7.4 (dd, J = 8.5 Hz and 2.5 Hz, 1H); 7.5 (d, J = 2.5 Hz, 1H); 7.6
15 (d, J = 16 Hz, 1H).

(d) Preparation of ethyl 3-(3',5'-di-tert-butyl-6-hydroxy-2'-pentyloxy-3-biphenyl)acrylate:

In a manner similar to that of Example 2(g),
starting with 1.84 g (3.6 mmol) of the above ethyl
20 ester, 1.6 g (88%) of the expected compound was obtained.

¹H NMR (d, CDCl₃): 0.85 (t, J = 7 Hz, 3H); 0.9-1.3 (m, 7H); 1.35 (s, 9H); 1.45 (s, 9H); 1.45-1.6 (m, 2H); 3.5-3.65 (m, 2H); 4.3 (q, J = 7 Hz, 2H); 6.35 (d, J = 16 Hz,
25 1H); 7.05 (d, J = 8 Hz, 1H); 7.15 (d, J = 2.4 Hz, 1H);

7.4 (d, J = 2.4 Hz, 1H); 7.5-7.6 (m, 2H); 7.7 (d, J = 16 Hz, 1H).

(e) Synthesis of 3-(3',5'-Di-tert-butyl-6-hydroxy-2'-pentyloxy-3-biphenyl)acrylic acid:

5 In a manner similar to that of Example 2(h), starting with 1.62 g (3.2 mmol) of ethyl 3-(3',5'-di-tert-butyl-6-hydroxy-2'-pentyloxy-3-biphenyl)acrylate, 1.2 g (86%) of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-pentyloxy-3-biphenyl)acrylic acid having a melting
10 point of 166°-7°C was obtained.
¹H NMR (d, CDCl₃): 0.8 (t, J = 7 Hz, 3H); 1.1-1.25 (m, 4H); 1.35 (s, 9H); 1.45 (s, 9H); 1.6 (t, J = 7 Hz, 2H); 3.5-3.6 (m, 2H); 6.4 (d, J = 16 Hz, 1H); 7.1 (d, J = 8 Hz, 1H); 7.2 (s, 1H); 7.4 (m, 2H); 7.5-7.6 (m, 2H); 7.8
15 (d, J = 16 Hz, 1H).

EXAMPLE 4:

Synthesis of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-methoxy-3-biphenyl)acrylic acid:

(a) Preparation of ethyl 3-(3',5'-di-tert-butyl-2'-methoxy-6-methoxymethoxy-3-biphenyl)acrylate:
20

In a manner similar to that of Example 1(c), by reaction of 2.18 g (8.26 mmol) of 3,5-di-tert-butyl-2-pentyloxyphenylboronic acid with 2 g (6.35 mmol) of

ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate, 2 g (70%) of the expected ethyl ester were obtained.

¹H NMR (d, CDCl₃): 1.2-1.3 (m, 3H); 1.35 (s, 9H); 1.45 (s, 9H); 3.3 (s, 3H); 3.4 (s, 3H); 3.8 (q, J = 7 Hz, 2H); 5.15 (s, 2H); 6.35 (d, J = 16 Hz, 1H); 7.1 (d, J = 2 Hz, 1H); 7.2 (d, J = 8 Hz, 1H); 7.35 (d, J = 2 Hz, 1H); 7.5 (dd, J = 8 Hz and J = 2 Hz, 1H); 7.6 (d, J = 2 Hz, 1H); 7.7 (d, J = 16 Hz, 1H).

10 (b) Preparation of ethyl 3-(3',5'-di-tert-butyl-6-hydroxy-2'-methoxy-3-biphenyl)acrylate:

In a manner similar to that of Example 2(g), starting with 1.45 g (3.17 mmol) of the above ethyl ester, 1.25 g (96%) of the expected compound was obtained.

15 ¹H NMR (d, CDCl₃): 1.1-1.2 (m, 4H); 1.35 (s, 9H); 1.45 (s, 9H); 3.5 (s, 3H); 4.25 (q, J = 7 Hz, 2H); 6.35 (d, J = 16 Hz, 1H); 7.05 (d, J = 8 Hz, 1H); 7.15 (d, J = 2.3 Hz, 1H); 7.4 (d, J = 2.3 Hz, 1H); 7.5-7.6 (m, 2H); 7.7 (d, J = 16 Hz, 1H).

20 (c) Synthesis of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-methoxy-3-biphenyl)acrylic acid:

In a manner similar to that of Example 2(h), starting with 1.23 g (3 mmol) of ethyl 3-(3',5'-di-tert-butyl-6-hydroxy-2'-methoxy-3-biphenyl)acrylate, 800 mg 25 (69%) of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-methoxy-3-

biphenylyl)acrylic acid having a melting point of 166°-7°C were obtained.

¹H NMR (d, CDCl₃): 1.35 (s, 9H); 1.45 (s, 9H); 3.5 (s, 3H); 6.4 (d, J = 16 Hz, 1H); 7.1 (d, J = 8 Hz, 1H); 7.2
5 (d, J = 2.5 Hz, 1H); 7.45 (d, J = 2.5 Hz, 1H); 7.5-7.6 (m, 2H); 7.8 (d, J = 16 Hz, 1H).

EXAMPLE 5:

Synthesis of 3-[3-(3-Benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid:
10

(a) Preparation of 3-benzyloxy-2-bromo-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a), by reaction of 5 g (25 mmol) of 3-hydroxy-2-bromo-
15 5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene with 3.5 ml (29.5 mmol) of benzyl bromide, 6.15 g (66%) of the expected compound were obtained.

¹H NMR (CDCl₃): 1.20 (s, 6H); 1.24 (s, 6H); 1.64 (s, 4H); 5.12 (s, 2H); 6.82 (s, 1H Ar); 7.21 to 7.50 (m, 6H
20 Ar).

(b) Preparation of 3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid:

In a manner similar to that of Example 1(b),
by reaction of 6.15 g (16.5 mmol) of 3-benzyloxy-2-
5 bromo-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene
with 5.7 ml (24.7 mmol) of triisopropyl borate, 3.5 g
(62%) of the expected boronic acid having a melting
point of 125°-6°C were obtained.

¹H NMR (CDCl₃): 1.26 (s, 6H); 1.29 (s, 6H); 1.68 (s,
10 4H); 5.12 (s, 2H); 5.83 (s, 2H); 6.88 (s, 1H Ar); 7.35
to 7.43 (m, 5H Ar); 7.80 (s, 1H Ar).

(c) Preparation of ethyl 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxy-phenyl]acrylate:

15 In a manner similar to that of Example 1(c),
by reaction of 3.4 g (10 mmol) of 3-benzyloxy-5,5,8,8-
tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid
with 3.17 g (10 mmol) of ethyl 3-(3-bromo-4-
methoxymethoxy-phenyl)acrylate, 3 g (56%) of the
20 expected ethyl ester were obtained.

¹H NMR (CDCl₃): 1.27 (s, 12H); 1.33 (t, J = 7.1 Hz, 3H);
1.69 (s, 4H); 3.31 (s, 3H); 4.25 (q, J = 7.1 Hz, 2H);
5.01 (s, 2H); 5.06 (s, 2H); 6.32 (d, J = 15.9 Hz, 1H);
6.89 (s, 1H Ar); 7.19 (s, 1H Ar); 7.21 to 7.31 (m, 6H
25 Ar), 7.46 (dd, J = 8.6 Hz, J = 2.3 Hz, 1H Ar); 7.52 (d,
J = 2.3 Hz, 1H Ar); 7.567 (d, J = 15.9 Hz, 1H).

(d) Synthesis of 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxy-phenyl]acrylic acid:

In a manner similar to that of Example 2(h), starting with 1.3 g (2.46 mmol) of the above ethyl ester, 1 g (90%) of 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid having a melting point of 161°-3°C was obtained.

¹H NMR (CDCl₃): 1.27 to 1.28 (m, 12H); 1.69 (s, 4H); 3.32 (s, 3H); 5.01 (s, 2H); 5.07 (s, 2H); 6.33 (d, J = 16 Hz, 1H); 6.70 (s, 1H Ar); 7.20 (s, 1H Ar); 7.21 to 7.31 (m, 6H Ar); 7.48 (dd, J = 8.6 Hz, J = 2.1 Hz, 1H Ar); 7.55 (d, J = 2 Hz, 1H Ar); 7.55 (d, J = 2 Hz, 1H).

¹³C NMR (CDCl₃): 1099, 32.11, 33.91, 34.69, 35.33, 56.20, 70.76, 95.02, 111.13, 115.23, 115.36, 125.11, 125.76, 127.08, 127.61, 127.68, 128.49, 129.25, 129.76, 129.83, 132.29, 137.37, 137.72, 145.74, 147.05, 153.85, 157.49, 172.54.

EXAMPLE 6:

Synthesis of 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-hydroxyphenyl]acrylic acid:

(a) Preparation of ethyl 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-hydroxyphenyl]acrylate:

In a manner similar to that of Example 2(g),
5 starting with 1.8 g (3.4 mmol) of ethyl 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylate, 1.2 g (72%)
of ethyl 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-hydroxyphenyl]acrylate was
10 obtained.

^1H NMR (CDCl_3): 1.25 to 1.29 (m, 12H); 1.31 (t, $J = 7.1$ Hz, 3H); 1.70 (s, 4H); 4.26 (q, $J = 7.1$ Hz, 2H); 5.07 (s, 2H); 6.33 (d, $J = 16$ Hz, 1H); 6.81 (s, 1H OH); 6.99 (s, 1H Ar); 7.01 (d, $J = 9.8$ Hz, 1H Ar); 7.23 (s, 1H
15 Ar); 7.26 to 7.32 (m, 5H Ar); 7.42 (d, $J = 2.1$ Hz, 1H Ar); 7.49 (dd, $J = 8.3$ Hz, $J = 2.1$ Hz, 1H Ar); 7.69 (d, $J = 16$ Hz, 1H).

(b) Synthesis of 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-hydroxyphenyl]acrylic acid:
20

In a manner similar to that of Example 2(h),
starting with 1.2 g (2.47 mmol) of the above ethyl ester, 980 mg (98%) of 3-[3-(3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-
25 4-hydroxyphenyl]acrylic acid having a melting point of $123^\circ\text{--}4^\circ\text{C}$ were obtained.

¹H NMR (CDCl₃): 1.28 to 1.30 (m, 12H); 1.70 (s, 4H);
5.08 (s, 2H); 6.35 (d, J = 15.7 Hz, 1H); 6.38 (s, 1H);
7.00 (s, 1H Ar); 7.04 (d, J = 7.5 Hz, 1H Ar); 7.24 (s,
1H Ar); 7.25 to 7.37 (m, 5H Ar); 7.45 (d, J = 2 Hz, 1H
5 Ar); 7.52 (dd, J = 8.4 Hz, J = 2 Hz, 1H Ar); 7.80 (d, J
= 15.9 Hz, 1H).

¹³C NMR (CDCl₃): 31.88, 32.02, 34.00, 34.65, 34.99,
35.08, 72.30, 112.66, 114.70, 118.16, 124.49, 127.07,
127.23, 127.64, 128.30, 128.68, 129.22, 130.55, 132.19,
10 136.06, 139.95, 147.01, 147.08, 152.48, 156.45, 172.75.

EXAMPLE 7:

Synthesis of [4-methoxymethoxy-3-(3,5,5,8,8-pentamethyl-
5,6,7,8-tetrahydro-2-naphthyl)phenyl]propynoic acid:

(a) Preparation of 2',2'-dibromo-4-
15 methoxymethoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-
tetrahydro-2-naphthyl)phenyl]ethylene:

4.95 g (14.71 mmol) of 4-methoxymethoxy-3-
(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)-
phenyl]carboxaldehyde and 50 ml of dichloromethane were
20 introduced into a round-bottomed flask. 9.75 g
(29.4 mmol) of carbon tetrabromide and 15.45 g
(58.84 mmol) of triphenylphosphine were added in
succession and the mixture was stirred at room
temperature for two hours. The reaction medium was
25 evaporated and the residue obtained was purified by

chromatography on a silica column eluted with dichloromethane. 6.75 g (88%) of the expected compound were collected.

5 (b) Preparation of [4-methoxymethoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acetylene:

7.6 g (12.9 mmol) of 2',2'-dibromo-4-methoxymethoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]ethylene and 50 ml of THF
10 were introduced under a stream of nitrogen into a three-necked flask. 11 ml (27.1 mmol) of a solution of n-butyllithium (2.5M in hexane) were added dropwise at -78°C and the mixture was permitted to cool to room temperature. The reaction medium was poured into water
15 and extracted with ethyl ether and the organic phase was decanted, dried over magnesium sulfate and evaporated. The residue obtained was purified by chromatography on a silica column eluted with a mixture of heptane and ethyl acetate (98:2). After evaporation of the solvents,
20 3.68 g (79%) of the expected acetylenic derivative were collected in the form of a yellow oil.

(c) Preparation of ethyl [4-methoxymethoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]propynoate:

3.65 g (10.1 mmol) of the above acetylenic
5 compound and 60 ml of THF were introduced under a stream
of nitrogen into a three-necked flask. 4.45 ml
(11.1 mmol) of n-butyllithium (2.5M in hexane) were
added dropwise at -78°C and the mixture was stirred for
thirty minutes. 1.15 ml (12.1 mmol) of ethyl
10 chloroformate was added at -78°C and the mixture was
permitted to cool to room temperature. The reaction
medium was poured into an aqueous ammonium chloride
solution and extracted with ethyl ether and the organic
phase was decanted, dried over magnesium sulfate and
15 evaporated. The residue obtained was purified by
chromatography on a silica column eluted with a mixture
of heptane and ethyl acetate (98:2). After evaporation
of the solvents, 4.14 g (95%) of ethyl [4-methoxy-
methoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-
20 naphthyl)phenyl]propynoate were collected in the form of
a yellow oil.

(d) Synthesis of [4-methoxymethoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]propynoic acid:

25 In a manner similar to that of Example 2(h),
starting from 439 mg. (1 mmol) of ethyl [4-methoxy-

methoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]propynoate, 375 mg (91%) of [4-methoxy-methoxy-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]propynoic acid having a melting point of
5 168°-170°C were obtained.

EXAMPLE 8:

Synthesis of 3-[3-(3-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid:

10 (a) **Preparation of ethyl 3-[3-(3-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylate:**

In a manner similar to that of Example 1(c),
by reaction of 3 g (10.4 mmol) of (3-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)boronic acid
15 (preparation described in WO 97/33881) with 3.2 g (10.1 mmol) of ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate (prepared in Example 2(c)), after
chromatography on a silica column and elution with a
20 mixture of heptane and ethyl acetate (98:2), 2.1 g (43%)
of the expected ethyl ester were obtained.

(b) Preparation of 3-[3-(3-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 724 mg (1.5 mmol) of ethyl 3-[3-(3-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylate, 620 mg (91%)
of 3-[3-(3-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic
10 acid having a melting point of 170°-1°C were obtained.

EXAMPLE 9:

Synthesis of 3-[3-(3-pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid:

15 (a) Preparation of 2-bromo-3-pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a),
by reaction of 5 g (25 mmol) of 3-bromo-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthol with 3.9 ml
20 (29.5 mmol) of 5-iodopentane, 7 g (79%) of the expected compound were obtained in the form of a colorless oil.

(b) Preparation of (3-pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)boronic acid:

In a manner similar to that of Example 1(b), starting from 6.9 g (19.5 mmol) of 2-bromo-3-pentyloxy-
5 5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene, 4.68 g (75%) of the expected boronic compound were obtained.

(c) Preparation of ethyl 3-[3-(3-pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylate:
10

In a manner similar to that of Example 1(c), by reaction of 4 g (12.6 mmol) of (3-pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)boronic acid with 2.64 g (8.4 mmol) of ethyl 3-[3-(3-bromo-4-methoxymethoxyphenyl)acrylate (prepared in
15 Example 2(c)), after chromatography on a silica column eluted with a mixture of heptane and ethyl acetate (98:2), 3.2 g (75%) of the expected ethyl ester were obtained.

(d) Synthesis of 3-[3-(3-pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid:
20

In a manner similar to that of Example 2(h), starting from 1.7 g (3.26 mmol) of ethyl 3-[3-(3-

5 pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylate, 1.35 g (86%) of 3-[3-(3-pentyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxymethoxyphenyl]acrylic acid having a melting point of 178°-80°C was obtained.

EXAMPLE 10:

Synthesis of 3-(5'-adamantan-1-yl-4'-methoxy-2'-methyl-3-biphenyl)acrylic acid:

(a) Preparation of 4-bromo-3-methylphenol:

10 80.0 g (740 mmol) of 3-methylphenol and 400 ml of glacial acetic acid were introduced under a nitrogen atmosphere into a 2 liter three-necked flask. The mixture was cooled to 15°C, 38 ml (742 mmol) of bromine were added dropwise, and the mixture was stirred at 15°C
15 for three hours. The reaction medium was poured into 1 liter of water and extracted with ethyl ether, the organic phase was decanted, washed with water to neutral pH and dried over magnesium sulfate, and the solvents were evaporated. The residue obtained was triturated in
20 heptane, filtered and dried. 70 g (50%) of a white powder having a melting point of 55°-56°C were collected.

(b) Preparation of 2-(1-adamantyl)-4-bromo-3-methylphenol:

70.0 g (376 mmol) of 4-bromo-3-methylphenol, 63 g (414 mmol) of 1-adamantanol, 500 ml of heptane and 50 ml of dichloromethane were introduced under a nitrogen atmosphere into a 2,000 ml round-bottomed flask. 20 ml (376 mmol) of 98% sulfuric acid were added dropwise and the mixture was stirred at room temperature for 3 hours. The reaction medium was poured into water, neutralized to a pH of 8-9 with sodium bicarbonate solution, and extracted with ethyl acetate, the organic phase was decanted, washed with water to neutral pH and dried over magnesium sulfate, and the solvents were evaporated. The residue obtained was triturated in heptane, filtered and dried. 59 g (49%) of a pinky white powder having a melting point of 139°-141°C were collected.

(c) Preparation of 2-(1-adamantyl)-4-bromo-5-methylanisole:

8 g (24.9 mmol) of 2-(1-adamantyl)-4-bromo-3-methylphenol, 75 ml of DMF and 25 ml of THF were introduced under a stream of nitrogen into a 250 ml three-necked flask. 880 mg (27.4 mmol) of sodium hydride (75% in oil) were added in small portions and the mixture was stirred until the evolution of gas had ceased. Then 1.9 ml (30 mmol) of iodomethane was added

and the mixture was stirred at room temperature for two hours. The reaction medium was poured into ice-water and extracted with ethyl ether and the organic phase was decanted, dried over magnesium sulfate and evaporated.

5 The residue obtained was triturated in heptane, filtered and dried. 6.44 g (77%) of pale-beige powder having a melting point of 129°-131°C were collected.

(d) Preparation of 5-(1-adamantyl)-4-methoxy-2-methylphenylboronic acid:

10 6 g (17.9 mmol) of 2-(1-adamantyl)-4-bromo-5-methylanisole and 40 ml of THF were introduced under a stream of nitrogen into a 250 ml three-necked flask. 7.9 ml (19.7 mmol) of n-butyllithium (2.5M in hexane) were added dropwise at -78°C and the mixture was stirred
15 for 15 minutes; at this same temperature, 5 ml (21.5 mmol) of triisopropyl borate were added and the mixture was stirred for 30 minutes. 20 ml of hydrochloric acid (1N) were added at -50°C and the mixture was permitted to warm to room temperature. The
20 reaction medium was extracted with ethyl ether and the organic phase was decanted, dried over magnesium sulfate and evaporated. The solid obtained was triturated in heptane, filtered and dried. 4.75 g (88%) of the expected boronic acid having a melting point of 276°-
25 278°C were collected, which was used as is in the remainder of the synthesis to follow.

(e) Synthesis of 3-(5'-adamantan-1-yl-4'-methoxy-2'-methyl-3-biphenyl)acrylic acid:

1.0 g (3.33 mmol) of 5-(1-adamantyl)-4-methoxy-2-methylphenylboronic acid obtained above,
5 580 mg (2.56 mmol) of 3-bromocinnamic acid and 30 ml of DME were introduced under a stream of nitrogen into a 100 ml three-necked flask. 4.5 ml of an aqueous potassium carbonate solution (2M) was added dropwise and the reaction medium was degassed. Then 150 mg
10 (0.13 mmol) of tetrakis(triphenylphosphine)palladium(0) were added and the mixture was heated at reflux for 2 hours. Water and ethyl acetate were added to the reaction medium, which was acidified to a pH of 1 with hydrochloric acid (1N). The organic phase was decanted,
15 washed with water, dried over magnesium sulfate and evaporated. The residue obtained was purified by chromatography on a silica column eluted with a mixture of heptane and ethyl acetate (85:15). After evaporation of the solvents, 750 mg of a thick gel were collected,
20 which were left to crystallize under a stream of nitrogen. The yellow crystals thus obtained were recrystallized from ethanol, filtered and dried to yield 452 mg (44%) of 3-(5'-adamantan-1-yl-4'-methoxy-2'-methyl-3-biphenyl)acrylic acid having a melting point of
25 194°-6°C.

¹H NMR (CDCl₃): 1.76 (br, 6H), 2.05 (br, 3H), 2.10 (br, 6H), 2.26 (s, 3H), 3.88 (s, 3H), 6.48 (d, J = 15.8 Hz,

1H), 6.78 (s, 1H Ar), 7.07 (s, 1H Ar), 7.35 to 7.51 (m, 4H Ar), 7.84 (d, J = 15.8 Hz, 1H Ar).

EXAMPLE 11:

5 **Synthesis of 3-(5'-adamantan-1-yl-6-hydroxy-4'-methoxy-2'-methyl-3-biphenyl)acrylic acid:**

(a) Preparation of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate:

10 In a manner similar to that of Example 2(g), by reaction of 10.1 g (20.3 mmol) of ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate obtained in Example 2(c) with 5.4 ml (102 mmol) of 98% sulfuric acid, 3.9 g of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate (49%) were obtained in the form of a white powder having a melting point of 112°-3°C.

15 **(b) Preparation of ethyl 3-(5'-adamantan-1-yl-6-hydroxy-4'-methoxy-2'-methyl-3-biphenyl)acrylate:**

20 In a manner similar to that of Example 1(c), by reaction of 3.0 g (10 mmol) of 5-(1-adamantyl)-4-methoxy-2-methylphenylboronic acid obtained in Example 10(d) with 2.25 g (8.33 mmol) of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate obtained above, 570 mg (15%) of a white powder having a melting point of 164°-6°C were obtained.

(c) Synthesis of 3-(5'-adamantan-1-yl-6-hydroxy-4'-methoxy-2'-methyl-3-biphenyl)acrylic acid:

In a manner similar to that of Example 2(h), starting from 270 mg (0.6 mmol) of ethyl 3-(5'-
5 adamantan-1-yl-6-hydroxy-4'-methoxy-2'-methyl-3-biphenyl)acrylate obtained above, 190 mg (76%) of a white powder which decomposed commencing at 120°C were obtained.

¹H NMR (CDCl₃): 1.76 (br, 6H), 2.07 (br, 9H), 2.13 (s, 3H), 3.89 (s, 3H), 5.20 (br, 1H), 6.31 (d, J = 15.75 Hz, 1H), 6.83 (s, 1H), 7.00 (d, J = 9.25 Hz, 1H), 7.48 (dd, J1 = 9.25 Hz, J2 = 2.0 Hz, 1H), 7.74 (d, J = 15.75 Hz, 1H).

EXAMPLE 12:

15 Synthesis of 3-(5'-adamantan-1-yl-4'-methoxy-6-methoxymethoxy-2'-methyl-3-biphenyl)acrylic acid:

(a) Preparation of ethyl 3-(5'-adamantan-1-yl-4'-methoxy-6-methoxymethoxy-2'-methyl-3-biphenyl)acrylate:

20 In a manner similar to that of Example 2(b), by reaction of 300 mg (0.67 mmol) of ethyl 3-(5'-adamantan-1-yl-6-hydroxy-4'-methoxy-2'-methyl-3-biphenyl)acrylate obtained in Example 11(b) with 64 mg

(0.8 mmol) of methoxymethyl chloride, 320 mg (97%) of a white gel were obtained.

¹H NMR (CDCl₃): 1.32 (t, J = 7.00 Hz, 3H), 1.75 (br, 6H), 2.04+2.09 (br+br, 6H+3H), 2.14 (s, 3H), 3.37 (s, 3H), 3.87 (s, 3H), 4.25 (q, J = 7.0 Hz, 2H), 5.11 (s, 2H), 6.33 (d, J = 15.8 Hz, 1H), 6.76 (s, 1H), 7.00 (s, 1H), 7.20 (d, J = 8.5 Hz, 1H), 7.36 (d, J = 2.25 Hz, 1H), 7.46 (dd, J₁ = 8.5 Hz, J₂ = 2.25 Hz, 1H), 7.66 (d, J = 15.8 Hz, 1H).

10 (b) Synthesis of 3-(5'-adamantan-1-yl-4'-methoxy-6-methoxymethoxy-2'-methyl-3-biphenyl)acrylic acid:

In a manner similar to that of Example 2(h), starting from 300 mg (0.61 mmol) of ethyl 3-(5'-adamantan-1-yl-4'-methoxy-6-methoxymethoxy-2'-methyl-3-biphenyl)acrylate obtained above, 200 mg (71%) of a white powder decomposed commencing at 113°C were obtained.

20 ¹H NMR (CDCl₃): 1.76 (br, 6H), 2.04 (br, 3H), 2.09 (br, 6H), 2.14 (s, 3H), 3.38 (s, 3H), 3.87 (s, 3H), 5.13 (s, 2H), 6.34 (d, J = 15.75 Hz, 1H), 6.76 (s, 1H), 7.00 (s, 1H), 7.22 (d, J = 8.5 Hz, 1H), 7.39 (d, J = 2.25 Hz, 1H), 7.49 (dd, J₁ = 8.5 Hz, J₂ = 2.25 Hz, 1H), 7.76 (d, J = 15.75 Hz, 1H).

EXAMPLE 13:

Synthesis of 3-{4-methoxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

5 (a) Preparation of 2-bromo-3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a), by reaction of 100 g (353 mmol) of 3-bromo-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthol with 32 ml
10 (406 mmol) of methoxymethyl chloride, 100 g (86%) of the expected compound were obtained in the form of a beige powder having a melting point of 69°-71°C.

(b) Preparation of 3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid:

15 In a manner similar to that of Example 1(b), by reaction of 90 g (275 mmol) of 2-bromo-3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-naphthalene obtained above, 70 g (87%) of the expected
20 compound were obtained in the form of a white powder having a melting point of 130°-132°C.

(c) Preparation of 3-bromo-4-methoxybenzaldehyde:

In a manner similar to that of Example 1(a),
by reaction of 50 g (270 mmol) of 3-bromo-4-
5 hydroxybenzaldehyde (obtained in accordance with
Example 2(a)) and 20.2 ml (338 mmol) of iodomethane,
48 g (83%) of the expected compound were obtained in the
form of a white powder having a melting point of
48°-50°C.

10 (d) Preparation of ethyl 3-(3-bromo-4-methoxyphenyl)acrylate:

In a manner similar to that of Example 2(c),
by reaction of 46 g (214 mmol) of 3-bromo-4-
methoxybenzaldehyde obtained above and 55.2 ml
15 (278 mmol) of triethyl phosphonoacetate, 47 g (77%) of
the expected compound were obtained in the form of white
flakes having a melting point of 76°-78°C.

20 (e) Preparation of ethyl 3-[4-methoxy-3-(3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylate:

In a manner similar to that of Example 1(c),
by reaction of 53.3 g (182 mmol) of 3-methoxymethoxy-
5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic
acid obtained in Example 13(b) and 40 g (140 mmol) of

ethyl 3-(3-bromo-4-methoxyphenyl)acrylate obtained above, 63 g (99%) of the expected compound were obtained in the form of a thick yellow oil.

¹H NMR (CDCl₃): 1.23 to 1.35 (m, 15H), 1.70 (s, 4H),
5 3.34 (s, 3H), 3.81 (s, 3H), 4.25 (q, J = 7.25 Hz, 2H),
5.03 (s, 2H), 6.32 (d, J = 15.75 Hz, 1H), 9.95 (d,
J = 9.0 Hz, 1H), 7.13 (s, 1H), 7.42 to 7.66 (m, 2H),
7.67 (d, J = 15.75 Hz, 1H).

(f) Preparation of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl, 5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate:

In a manner similar to that of Example 2(g), by reaction of 63 g (140 mmol) of ethyl 3-[4-methoxy-3-(3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylate obtained above and
15 22 ml (420 mmol) of sulfuric acid, 42.5 g (75%) of the expected compound were obtained in the form of a pale-yellow powder with a melting point of 147-149°C.

(g) Preparation of ethyl 3-{4-methoxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 1(a), by reaction of 1 g (2.45 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained above with 4 ml (3 mmol)
25

of 3-methoxybenzyl chloride, 1.30 g (100%) of the expected compound was obtained in the form of a yellow oil.

¹H NMR (CDCl₃): 1.24 to 1.35 (m, 15H), 1.69 (s, 4H),
5 3.69 (s, 3H), 3.76 (s, 3H), 4.25 (q, J = 7.0 Hz, 2H),
5.0 (s, 2H), 6.30 (d, J = 16.0 Hz, 1H), 6.75 to 6.96 (m, 5H), 7.16 to 7.22 (m, 2H), 7.45 to 7.51 (m, 2H), 7.67 (d, J = 16.0 Hz, 1H).

(h) Synthesis of 3-{4-methoxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 1.3 g (2.45 mmol) of ethyl 3-{4-methoxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above,
15 830 mg (67%) of the expected compound were obtained in the form of a white powder having a melting point of 168°-170°C.

EXAMPLE 14:

20 Synthesis of 3-{4-methoxy-3-[3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

(a) Preparation of ethyl 3-{4-methoxy-3-[3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 1(a),
5 by reaction of 1.23 g (3 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f) with 490 μ l (3.6 mmol) of 4-methoxybenzyl chloride, 1.58 g (100%) of the expected product are obtained in the form
10 of a yellow oil.

^1H NMR (CDCl_3): 1.21 to 1.35 (m, 15H), 1.69 (s, 4H),
3.75 (s, 3H), 3.77 (s, 3H), 4.25 (q, $J = 7.0$ Hz, 2H),
4.93 (s, 2H), 6.29 (d, $J = 16.0$ Hz, 1H), 6.80 to 6.94
(m, 4H), 7.15 to 7.18 (m, 2H), 7.31 (d, $J = 8.5$ Hz, 1H),
15 7.44 to 7.50 (m, 2H), 7.66 (d, $J = 16.25$ Hz, 1H).

(b) Synthesis of 3-{4-methoxy-3-[3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h),
20 starting from 1.58 g (3 mmol) of ethyl 3-{4-methoxy-3-[3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 1.1 g (71%) of the expected compound was obtained in the form of a white powder having a melting point of 201° -
25 203°C .

EXAMPLE 15:

Synthesis of 3-{3-[3-(6-hydroxyhexyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

- 5 (a) Preparation of ethyl 3-{3-[3-(6-hydroxyhexyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

In a manner similar to that of Example 1(a), by reaction of 1.4 g (3.4 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f) with 540 μ l (4.1 mmol) of 1-bromohexanol, 850 mg (50%) of the expected compound were obtained in the form of a yellow oil.

15 ^1H NMR (CDCl_3): 1.23 to 1.35 (m, 19H), 1.43 to 1.70 (m, 8H), 3.56 (q, $J = 5.5$ Hz, 2H), 3.81 (s, 3H), 3.90 (t, $J = 5.0$ Hz, 2H), 4.25 (q, $J = 7.25$ Hz, 2H), 6.31 (d, $J = 16.0$ Hz, 1H), 6.85 (s, 1H), 6.96 (d, $J = 8.5$ Hz, 1H), 7.20 (s, 1H), 7.47 (dd, $J_1 = 8.25$ Hz, $J_2 = 2.25$ Hz, 1H), 7.52 (d, $J = 2.25$ Hz, 1H), 7.67 (d, $J = 16.0$ Hz, 1H).

20

(b) Synthesis of 3-{3-[3-(6-hydroxyhexyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 850 mg (1.67 mmol) of ethyl 3-{3-[3-(6-hydroxyhexyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained above,
510 mg (63%) of the expected compound were obtained in
the form of a white powder having a melting point of
10 118°-120°C.

EXAMPLE 16:

Synthesis of 3-{3-[3-(7-hydroxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

15 (a) Preparation of ethyl 3-{3-[3-(7-hydroxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

1.23 g (3 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f), 440 mg
20 (3.2 mmol) of potassium carbonate, 540 μ l (3.5 mmol) of 1-bromoheptanol and 30 ml of butanone were introduced into a round-bottomed flask. The mixture was heated at reflux for 8 hours, the reaction medium was poured into

water, acidified to a pH of 1 with hydrochloric acid and extracted with ethyl ether, and the organic phase was decanted, dried over magnesium sulfate and evaporated. 1.57 g (100%) of the expected compound in the form of a yellow oil was collected.

¹H NMR (CDCl₃): 1.26 (s, 6H), 1.32 (s, 6H), 1.08 to 1.7 (m, 17H), 3.5 to 3.7 (m, 2H), 3.81 (s, 3H), 3.89 (t, J = 6.25 Hz, 2H), 4.24 (q, J = 7.0 Hz, 2H), 6.3 (d, J = 16.0 Hz, 1H), 6.84 (s, 1H), 6.95 (d, J = 8.25 Hz, 1H), 7.19 (s, 1H), 7.47 (dd, J₁ = 8.25 Hz, J₂ = 2.25 Hz, 1H), 7.50 (d, J = 2.25 Hz, 1H), 7.67 (d, J = 16.0 Hz, 1H).

(b) Synthesis of 3-{3-[3-(7-hydroxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 1.57 g (3 mmol) of ethyl 3-{3-[3-(7-hydroxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained above, 700 mg (47%) of the expected compound were obtained in the form of a white powder having a melting point of 105°-107°C.

EXAMPLE 17:

Synthesis of 3-{3-[3-(5-hydroxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

- 5 (a) Preparation of ethyl 3-{3-[3-(5-acetoxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

In a manner similar to that of Example 1(a), by reaction of 1.20 g (2.90 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f) with 727 mg (3.48 mmol) of 5-bromopentyl acetate, 1.42 g (92%) of the expected compound was obtained in the form of a yellow oil.

15 ¹H NMR (CDCl₃): 1.26 to 1.37 (m, 17H), 1.50 to 1.67 (m, 4H), 1.70 (s, 4H), 2.01 (s, 3H), 3.80 (s, 3H), 3.80 to 4.00 (m, 4H), 4.25 (q, J = 7.25 Hz, 2H), 6.30 (d, J = 15.75 Hz, 1H), 6.84 (s, 1H), 6.94 (d, J = 8.25 Hz, 1H), 7.17 (s, 1H), 7.44 to 7.48 (m, 2H), 7.66 (d, 20 J = 16.0 Hz, 1H).

(b) Synthesis of 3-{3-[3-(5-hydroxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 1.42 g (2.65 mmol) of ethyl 3-{3-[3-(5-acetoxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained
above, 730 mg (60%) of the expected compound were
obtained in the form of a white powder having a melting
10 point of 115°-117°C.

EXAMPLE 18:

Synthesis of 3-{3-[3-(3-Hydroxypropyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

15 (a) Preparation of ethyl 3-{3-[3-(3-Hydroxypropyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

In a manner similar to that of Example 16(a),
by reaction of 1.2 g (2.9 mmol) of ethyl 3-[3-(3-
20 hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl]acrylate obtained in
Example 13(f) with 490 mg (3.5 mmol) of 1-bromopropanol,
632 mg (47%) of the expected compound were obtained in
the form of a yellow oil.

¹H NMR (CDCl₃): 1.27 to 1.35 (m, 9H), 1.70 (s, 4H), 1.85 to 1.92 (m, 2H), 1.96 (br, 1H), 3.64 (m, 2H), 3.82 (s, 3H), 4.09 (t, J = 5.5 Hz, 2H), 4.25 (q, J = 7.25 Hz, 2H), 6.31 (d, J = 16.0 Hz, 1H), 6.87 (s, 1H), 6.96 (d, J = 8.5 Hz, 1H), 7.15 (s, 1H), 7.44 (d, J = 2.25 Hz, 1H), 7.49 (dd, J₁ = 8.5 Hz, J₂ = 2.25 Hz, 1H), 7.66 (d, J = 16.0 Hz, 1H).

(b) Synthesis of 3-{3-[3-(3-hydroxypropyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 632 mg (1.35 mmol) of ethyl 3-{3-[3-(3-hydroxypropyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained above, 449 mg (75%) of the expected compound obtained in the form of a white powder having a melting point of 198°-200°C.

EXAMPLE 19:

Synthesis of the intermediate 3-[3-(1-methoxymethoxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

(a) Preparation of 1-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a),
by reaction of 8.65 g (42.4 mmol) of 5,5,8,8-
5 tetramethyl-5,6,7,8-tetrahydro-2-naphthol with 3.55 ml
(46.6 mmol) of methoxymethyl chloride, 10.5 g (100%) of
the expected compound were obtained.

(b) Preparation of 1-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthaleneboronic
10 acid:

1 g (4 mmol) of 1-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene and 12 ml of THF were introduced into a round-bottomed flask. The mixture was cooled to -78°C and 630 µl (4.2 mmol) of
15 TMEDA and then 3.4 ml (4.4 mmol) of sec-BuLi (1.3M) were added dropwise. The mixture was stirred at -78°C for 1 hour and then 1.4 ml (6 mmol) of triisopropyl borate was added and the mixture was permitted to warm to room temperature. 3 ml of hydrochloric acid (5%) were added
20 at -40°C and the mixture was stirred at room temperature for 1 hour. The reaction medium was poured into water and extracted with ethyl acetate and the organic phase was decanted, dried over magnesium sulfate and evaporated. The residue obtained is purified by
25 chromatography on a silica column eluted with a mixture of dichloromethane and methanol (98:2). 900 mg (77%) of

the expected boronic compound in the form of a pale-yellow oil were collected.

5 (c) Synthesis of 3-[3-(1-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid:

 In a manner similar to that of Example 1(c), by reaction of 3.4 g (11.6 mmol) of 1-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthalene-boronic acid with 1.32 g (5.8 mmol) of 3-bromocinnamic
10 acid, trituration in heptane gave 1.4 g (62%) of 3-[3-(1-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid in the form of a white powder having a melting point of 188°-9°C.

EXAMPLE 20:

15 Synthesis of 3-[3-(1-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid:

 (a) Preparation of methyl 3-[3-(1-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylate:

20 1.24 g (3.15 mmol) of 3-[3-(1-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid obtained in Example 19(c), 15 ml of methanol and 15 ml of THF were introduced into

a round-bottomed flask. 620 μ l of concentrated sulfuric acid were added and the mixture was stirred at room temperature for eight hours and heated at reflux for two hours. The reaction medium was poured into water and
5 extracted with ethyl ether and the organic phase was decanted, dried over magnesium sulfate and evaporated. 1.1 g (100%) of methyl 3-[3-(1-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]-acrylate having a melting point of 197°-8°C were
10 collected.

(b) Preparation of methyl 3-[3-(1-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylate:

In a manner similar to that of Example 1(a),
15 by reaction of 500 mg (1.37 mmol) of methyl 3-[3-(1-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylate with 180 μ l (1.5 mmol) of benzyl bromide, 550 mg (88%) of the expected compound were obtained in the form of a yellow oil.

20 (c) Synthesis of 3-[3-(1-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid:

In a manner similar to that of Example 2(h),
starting from 498 mg (1.01 mmol) of ethyl 3-[3-(1-
25 benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-

naphthyl)phenyl]acrylate, 360 mg (82%) of 3-[3-(1-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid having a melting point of 153°-4°C were obtained.

5

EXAMPLE 21:

Synthesis of 3-{3-[3-(5-tert-butoxycarbonylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

10 (a) Preparation of tert-butyl 6-bromohexanoate:

1.95 g (10 mmol) of 6-bromohexanoic acid dissolved in 10 ml of dichloromethane were introduced under a stream of nitrogen into a round-bottomed flask. A solution of 4.37 g (20 mmol) of tert-butyl 2,2,2-trichloroacetimidate in 20 ml of cyclohexane was added dropwise, followed immediately by 200 μ l of trifluoroborane etherate. The mixture was stirred at room temperature for 5 minutes, 5 g of sodium bicarbonate were added, and the mixture was stirred for 1 minute. The suspension obtained was placed on a silica column which had been wetted beforehand with cyclohexane and the latter was eluted with a 8:2 cyclohexane/ethyl acetate mixture. After evaporation of the solvents, 1.63 g (65%) of a colorless oil was obtained.

^1H NMR (CDCl_3): 1.45 (s, 9H), 1.44 to 1.70 (m, 4H), 1.87 (m, 2H), 2.23 (t, $J = 7.5$ Hz, 2H), 3.41 (t, $J = 6.75$ Hz, 2H).

5 (b) Preparation of ethyl 3-{3-[3-(5-tert-butoxycarbonylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

In a manner similar to that of Example 1(a), by reaction of 2.36 g (5.8 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in
10 Example 13(f) with 1.6 g (6.3 mmol) of tert-butyl 6-bromohexanoate obtained above, 3.3 g (100%) of the expected compound were obtained in the form of a yellow oil.

15 ^1H NMR (CDCl_3): 1.21 to 1.75 (m, 32H), 2.13 (t, $J = 7.25$ Hz, 2H), 3.80 (s, 3H), 3.88 (t, $J = 6.25$ Hz, 2H), 4.25 (q, $J = 7.0$ Hz, 2H), 6.30 (d, $J = 16.0$ Hz, 1H), 6.83 (s, 1H), 6.94 (d, $J = 8.75$ Hz, 1H), 7.16 (s, 1H), 7.44 to 7.48 (m, 2H), 7.66 (d, $J = 16.0$ Hz, 1H).

20 (c) Synthesis of 3-{3-[3-(5-tert-butoxycarbonylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

A solution of 1.0 g (1.7 mmol) of ethyl 3-{3-[3-(5-tert-butoxycarbonylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-
25

methoxyphenyl}acrylate obtained above dissolved in 25 ml of THF was placed in a round-bottomed flask and 730 mg (17 mmol) of lithium hydroxide and 2 ml of water were added. The mixture was heated at the reflux of the THF for 8 hours and stirred at room temperature for 48 hours, the reaction medium was poured into water, acidified to a pH of 1 with hydrochloric acid and extracted with ethyl ether, and the organic phase was decanted, dried over magnesium sulfate, filtered and evaporated. The residue obtained was chromatographed on a silica column eluted with a 8:2 heptane/ethyl acetate mixture. After evaporation of the solvents, 126 mg (14%) of a white powder having a melting point of 130°-132°C were collected.

EXAMPLE 22:

Synthesis of 3-{3-[3-(7-tert-butoxycarbonylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

(a) Preparation of tert-butyl 8-bromooctanoate:

In a manner similar to that of Example 21(a), starting from 3 g (13.4 mmol) of 8-bromooctanoic acid, 3 g (80%) of a pale-yellow oil were obtained.

¹H NMR (CDCl₃): 1.29 to 1.37 (m, 6H), 1.44 (s, 9H), 1.50 to 1.75 (m, 2H), 1.75 to 1.95 (m, 2H), 2.20 (t, J = 7.5 Hz, 2H), 3.40 (t, J = 6.75 Hz, 2H).

(b) Preparation of ethyl 3-{3-[3-(7-tert-butoxycarbonylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

In a manner similar to that of Example 1(a),
5 by reaction of 3.82 g (9.34 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in
Example 13(f) with 3 g (10.7 mmol) of tert-butyl
8-bromooctanoate obtained above, 4.36 g (77%) of the
10 expected compound were obtained in the form of a yellow
oil.

¹H NMR (CDCl₃): 1.26 to 1.13 (m, 21H), 1.44 (s, 9H),
1.43 to 1.65 (m, 4H), 1.70 (s, 4H), 2.16 (t, J =
7.25 Hz, 2H), 3.80 (s, 3H), 3.88 (t, J = 6.5 Hz, 2H),
15 4.24 (q, J = 7.0 Hz, 2H), 6.30 (d, J = 15.75 Hz, 1H),
6.84 (s, 1H), 6.93 (d, J = 8.25 Hz, 1H), 7.17 (s, 1H),
7.43 to 7.49 (m, 2H), 7.66 (d, J = 15.75 Hz, 1H).

(c) Synthesis of 3-{3-[3-(7-tert-butoxycarbonylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 21(c),
starting from 1.15 g (1.9 mmol) of ethyl 3-{3-[3-(7-tert-butoxycarbonylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate
25 obtained above, 363 mg (33%) of the expected compound

were obtained in the form of a white powder having a melting point of 126°-128°C.

EXAMPLE 23:

Synthesis of 3-{3-[3-(7-Carboxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

(a) Preparation of ethyl 3-{3-[3-(7-carboxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

10 A solution of 3.75 g (6.18 mmol) of ethyl
3-{3-[3-(7-tert-butoxycarbonylheptyloxy)-5,5,8,8-
tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-
methoxyphenyl}acrylate obtained in Example 22(b)
dissolved in 60 ml of dichloromethane was placed in a
15 round-bottomed flask, 4.76 ml (61.8 mmol) of
trifluoroacetic acid were added, and the mixture was
stirred at room temperature for 4 hours. The reaction
medium was poured into water and extracted with ethyl
ether, and the organic phase was decanted, dried over
20 magnesium sulfate, filtered and evaporated.
3.4 g (100%) of the expected compound were obtained in
the form of a white foam.
¹H NMR (CDCl₃): 1.26 to 1.35 (m, 21H), 1.50 to 1.67 (m,
4H), 1.70 (s, 4H), 2.30 (t, J = 7.5 Hz, 2H), 3.80 (s,
25 3H), 3.88 (t, J = 6.25 Hz, 2H), 4.25 (q, J = 7.25 Hz,

2H), 6.31 (d, J = 15.75 Hz, 1H), 6.84 (s, 1H), 6.94 (d, J = 8.25 Hz, 1H), 7.17 (s, 1H), 7.44 to 7.49 (m, 2H), 7.67 (d, J = 16.0 Hz, 1H).

(b) Synthesis of 3-{3-[3-(7-carboxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 1 g (1.8 mmol) of ethyl 3-{3-[3-(7-carboxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained above, 830 mg (88%) of the expected compound were obtained in the form of a white powder having a melting point of 212°-214°C.

EXAMPLE 24:

Synthesis of 3-{3-[3-(5-carboxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

(a) Preparation of ethyl 3-{3-[3-(5-carboxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

In a manner similar to that of Example 23(a), starting from 2.3 g (3.97 mmol) of ethyl 3-{3-[3-(5-tert-butoxycarbonylpentyloxy)-5,5,8,8-tetramethyl-

5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained in Example 21(b), 2.1 g (100%) of the expected compound were obtained in the form of a white foam.

¹H NMR (CDCl₃): 1.24 to 1.42 (m, 17H), 1.55 to 1.70 (m, 8H), 2.28 (t, J = 7.0 Hz, 2H), 3.80 (s, 3H), 3.91 (t, J = 5.75 Hz, 2H), 4.25 (q, J = 7.25 Hz, 2H), 6.32 (d, J = 15.75 Hz, 1H), 6.84 (s, 1H), 6.95 (d, J = 9.0 Hz, 1H), 7.19 (s, 1H), 7.46 to 7.50 (m, 2H), 7.69 (d, J = 16.0 Hz, 1H).

10 (b) Synthesis of 3-{3-[3-(5-carboxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 900 mg (1.72 mmol) of ethyl 3-{3-[3-(5-carboxypentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained above, 647 mg (76%) of the expected compound were obtained in the form of a white powder having a melting point of 166°-168°C.

20 EXAMPLE 25:

Synthesis of 3-{3-[3-(5-carbamoylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

(a) Preparation of ethyl 3-{3-[3-(5-Carbamoylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

A solution of 1.27 g (2.43 mmol) of ethyl
5 3-{3-[3-(5-carboxypentyloxy)-5,5,8,8-tetramethyl-
5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate
obtained in Example 24(a) dissolved in 35 ml of
dichloromethane was introduced into a round-bottomed
flask, 505 μ l (2.55 mmol) of dicyclohexylamine were
10 added, and the mixture was stirred for 10 minutes. Then
186 μ l (2.55 mmol) of thionyl chloride were added
dropwise and the mixture was stirred at room temperature
for 10 minutes. The reaction medium was evaporated to
dryness, the residue obtained was taken up in ethyl
15 ether, the suspension was filtered and the filtrate was
evaporated. The residue obtained was dissolved in 15 ml
of THF and the solution was introduced dropwise under
nitrogen into a three-necked flask containing 20 ml of
THF, 405 μ l (2.92 mmol) of triethylamine and 160 μ l of a
20 34% aqueous ammonia solution (2.67 mmol). The mixture
was stirred at room temperature for 30 minutes, the
reaction medium was poured into water, acidified to a pH
of 1 with 1N hydrochloric acid and extracted with ethyl
ether, and the organic phase was decanted, dried over
25 magnesium sulfate, filtered and evaporated. The residue
obtained was purified by chromatography on a silica
column eluted with a mixture of heptane and ethyl

acetate (80:20). After evaporation of the solvents, 750 mg (60%) of the expected compound were obtained in the form of a white powder having a melting point of 133°-135°C.

5 (b) Synthesis of 3-{3-[3-(5-carbamoylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 900 mg (1.4 mmol) of ethyl 3-{3-[3-(5-carbamoylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained
10 above, 190 mg (27%) of the expected compound were obtained in the form of a white powder having a melting point of 179°-181°C.

15 EXAMPLE 26:

Synthesis of 3-{3-[3-(7-carbamoylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

20 (a) Preparation of ethyl 3-{3-[3-(7-carbamoylpentyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate:

In a manner similar to that of Example 25(a), starting from 1.5 g (2.70 mmol) of ethyl 3-{3-[3-(7-

carboxyheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained in Example 23(a), 1 g (68%) of the expected compound was obtained in the form of a beige powder having a melting point of 132°-134°C.

(b) Synthesis of 3-{3-[3-(7-carbamoylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 1 g (1.8 mmol) of ethyl 3-{3-[3-(7-carbamoylheptyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]-4-methoxyphenyl}acrylate obtained above, 700 mg (74%) of the expected compound were obtained in the form of a white powder having a melting point of 199°-201°C.

EXAMPLE 27:

Synthesis of 3-{4-methoxy-3-[5,5,8,8-tetramethyl-3-(2-morpholin-4-yl-ethoxy)-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

(a) Preparation of ethyl 3-{4-methoxy-3-[5,5,8,8-tetramethyl-3-(2-morpholin-4-yl-ethoxy)-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 16(a),
5 by reaction of 1.4 g (3.4 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f) with 760 mg (4.08 mmol) of 4-(2-chloroethyl)-morpholine hydrochloride, 1.5 g (85%) of the expected
10 compound was obtained in the form of a yellow oil.
¹H NMR (CDCl₃): 1.27 (s, 6H), 1.29 to 1.35 (m, 9H), 1.70 (s, 4H), 2.37 (m, 4H), 2.60 (t, J = 5.75 Hz, 2H), 3.59 (m, 4H), 3.81 (s, 3H), 4.05 (t, J = 5.50 Hz, 2H), 4.25 (q, J = 7.0 Hz, 2H), 6.30 (d, J = 15.75 Hz, 1H), 6.85
15 (s, 1H), 6.91 to 6.95 (m, 1H), 7.16 (s, 1H), 7.44 to 7.49 (m, 2H), 7.65 (d, J = 16.0 Hz, 1H).

(b) Synthesis of 3-{4-methoxy-3-[5,5,8,8-tetramethyl-3-(2-morpholin-4-yl-ethoxy)-5,6,7,8-tetrahydro-2-naphthyl]phenyl}-acrylic acid:

20 In a manner similar to that of Example 2(h), starting from 1.5 g (2.9 mmol) of ethyl 3-{4-methoxy-3-[5,5,8,8-tetramethyl-3-(2-morpholin-4-yl-ethoxy)-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 1.17 g (83%) of the expected compound was
25 obtained in the form of a white powder which decomposed commencing at 140°C.

¹H NMR (CD₃OD (Ref.: q at δ = 2.85 ppm)): 0.79 (s, 6H),
0.87 (s, 6H), 1.26 (s, 4H), 1.90 (m, 4H), 2.18 (t, J =
5.0 Hz, 2H), 3.07 (m, 4H), 3.31 (s, 3H), 3.61 (t, J =
5.0 Hz, 2H), 5.91 (d, J = 15.75 Hz, 1H), 6.45 (s, 1H),
5 6.56 (d, J = 8.5 Hz, 1H), 6.62 (s, 1H), 6.87 (d, J =
1.75 Hz, 1H), 7.01 to 7.09 (m, 2H).

EXAMPLE 28:

Synthesis of 3-{4-methoxy-3-[5,5,8,8-tetramethyl-3-(2-
piperidin-1-yl-ethoxy)-5,6,7,8-tetrahydro-2-
10 naphthyl]phenyl}acrylic acid:

(a) Preparation of ethyl 3-{4-methoxy-3-
[5,5,8,8-tetramethyl-3-(2-piperidin-1-yl-ethoxy)-
5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 16(a),
15 by reaction of 1.4 g (3.4 mmol) of ethyl 3-[3-(3-
hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-
naphthyl)-4-methoxyphenyl]acrylate obtained in Example
13(f) and 760 mg (4.08 mmol) of 1-(2-chloroethyl)-
piperidine hydrochloride, 1.57 g (89%) of the expected
20 compound was obtained in the form of a colorless oil.
¹H NMR (CDCl₃): 1.22 to 1.38 (m, 16H), 1.46 to 1.55 (m,
4H), 1.69 (m, 5H), 2.36 (m, 4H), 2.60 (t, J = 6.25 Hz,
2H), 3.81 (s, 3H), 4.05 (t, J = 6.0 Hz, 2H), 4.24 (q,
J = 7.25 Hz, 2H), 6.30 (d, J = 16.0 Hz, 1H), 6.85 (s,
25 1H), 6.93 (d, J = 9.25 Hz, 1H), 7.16 (s, 1H), 7.44 to
7.48 (m, 2H), 7.66 (d, J = 16.0 Hz, 1H).

(b) Synthesis of 3-{4-methoxy-3-[5,5,8,8-tetramethyl-3-(2-piperidin-1-yl-ethoxy)-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 1.57 g (3 mmol) of ethyl 3-{4-methoxy-3-[5,5,8,8-tetramethyl-3-(2-piperidin-1-yl-ethoxy)-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 900 mg (61%) of the expected compound were
obtained in the form of a white powder having a melting
10 point of 251°-253°C.

EXAMPLE 29:

Synthesis of 3-{4-methoxy-3-[3-(2-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

15 (a) Preparation of 2-methoxymethoxybenzaldehyde:

In a manner similar to that of Example 1(a),
by reaction of 30 g (245 mmol) of 2-hydroxybenzaldehyde
with 21 ml (270 mmol) of methoxymethyl chloride, 40.7 g
(100%) of the expected compound were obtained in the
20 form of an off-white oil.

¹H NMR (CDCl₃): 3.53 (s, 3H), 5.31 (s, 2H), 7.09 (t, J =
7.5 Hz, 1H), 7.22 (d, J = 8.25 Hz, 1H), 7.50 to 7.57 (m,
1H), 7.85 (dd, J₁ = 7.75 Hz, J₂ = 1.75 Hz, 1H), 10.51
(s, 1H).

(b) Preparation of (2-methoxymethoxyphenyl)methanol:

A solution of 40 g (241 mmol) of 2-methoxymethoxybenzaldehyde dissolved in 250 ml of THF was introduced under a stream of nitrogen into a 2-liter round-bottom flask, and 3.65 g (96 mmol) of lithium aluminum hydride were added in small portions. The reaction medium was stirred at room temperature for 15 minutes, before being hydrolyzed slowly with a saturated ammonium chloride solution and was then acidified to a pH of 1 with a 1N hydrochloric acid solution. The reaction medium was extracted with ethyl acetate, the organic phase was decanted, dried over magnesium sulfate and filtered, and the filtrate was evaporated. 39.2 g (96%) of a colorless oil were collected.

^1H NMR (CDCl_3): 2.29 (m, 1H), 3.49 (s, 3H), 4.80 (d, $J = 6.0$ Hz, 2H), 5.25 (s, 2H), 6.98 to 7.12 (m, 2H), 7.23 to 7.34 (m, 2H).

(c) Preparation of 2-methoxymethoxybenzyl chloride:

5.0 g (29.7 mmol) of (2-methoxymethoxyphenyl)-methanol obtained above, 90 ml of dichloromethane and 8.27 ml (59.5 mmol) of triethylamine were introduced in succession under a stream of nitrogen into a 250 ml three-necked flask. Then 2.76 g (35.7 mmol) of

methanesulfonyl chloride were added dropwise and the mixture was stirred at room temperature for 1 hour. The reaction medium was hydrolyzed with hydrochloric acid and extracted with dichloromethane, the organic phase
5 was dried over magnesium sulfate and filtered, and the filtrate was evaporated. The orange oil obtained was purified by chromatography on a silica column eluted with a mixture of ethyl acetate and heptane (1:9). After evaporation of the solvents, 3.0 g (55%) of the expected
10 compound in the form of a yellow oil were collected.
¹H NMR (CDCl₃): 3.50 (s, 3H), 4.67 (s, 2H), 3.25 (s, 2H), 6.99 (td, J₁ = 7.5 Hz, J₂ = 1.0 Hz, 1H), 7.11 (d, J = 7.75 Hz, 1H), 7.24 to 7.38 (m, 2H).

15 (d) Preparation of ethyl 3-{4-methoxy-3-[3-(2-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 1(a), by reaction of 2.5 g (6.12 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-
20 naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f) with 1.26 g (6.73 mmol) of 2-methoxymethoxybenzyl chloride obtained above, 3.2 g (94%) of the expected compound were obtained in the form of a yellow oil.
¹H NMR (CDCl₃): 1.27 to 1.35 (m, 15H), 1.69 (s, 4H),
25 3.43 (s, 3H), 3.74 (s, 3H), 4.25 (q, J = 7.0 Hz, 2H), 5.07 (s, 2H), 5.18 (s, 2H), 6.29 (d, J = 16.0 Hz, 1H),

6.90 to 6.95 (m, 3H), 7.06 (d, J = 7.5 Hz, 1H), 7.15 to 7.21 (m, 3H), 7.47 (dd, J₁ = 8.5 Hz, J₂ = 2.25 Hz, 1H), 7.52 (d, J = 2.25 Hz, 1H), 7.66 (d, J = 16.0 Hz, 1H).

5 (e) Synthesis of 3-{4-methoxy-3-[3-(2-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

10 In a manner similar to that of Example 2(h), starting from 1.0 g (1.8 mmol) of ethyl 3-{4-methoxy-3-[3-(2-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 760 mg (80%) of the expected compound were obtained in the form of a white powder having a melting point of 159°-161°C.

EXAMPLE 30:

15 Synthesis of 3-{4-methoxy-3-[3-(3-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

 (a) Preparation of 3-methoxymethoxybenzaldehyde:

20 In a manner similar to that of Example 1(a), by reaction of 30 g (245 mmol) of 3-hydroxybenzaldehyde with 21 ml (270 mmol) of methoxymethyl chloride, 40.7 g

(100%) of the expected compound were obtained in the form of a chestnut-brown oil.

^1H NMR (CDCl_3): 3.49 (s, 3H), 5.23 (s, 2H), 7.27 to 7.32 (m, 1H), 7.42 to 7.55 (m, 3H), 9.97 (s, 1H).

5 (b) Preparation of (3-methoxymethoxyphenyl)methanol:

In a manner similar to that of Example 29(b), by reaction of 40 g (240 mmol) of 3-methoxymethoxybenzaldehyde with 3.65 g (96 mmol) of lithium aluminum
10 hydride, 40.7 g (100%) of the expected compound were obtained in the form of a colorless oil.

^1H NMR (CDCl_3): 1.83 (br, 1H), 3.48 (s, 3H), 4.67 (s, 2H), 5.18 (s, 2H), 6.94 to 7.06 (m, 3H), 7.24 to 7.31 (m, 1H).

15 (c) Preparation of 3-methoxymethoxybenzyl chloride:

In a manner similar to that of Example 29(c), by reaction of 5.0 g (29.7 mmol) of
(3-methoxymethoxyphenyl)methanol with 8.27 ml
20 (59.5 mmol) of triethylamine and 2.76 ml (35.7 mmol) of methanesulfonyl chloride, 2.66 g (48%) of the expected compound were obtained in the form of a yellow oil.

^1H NMR (CDCl_3): 3.48 (s, 3H), 4.55 (s, 2H), 5.22 (s, 2H), 6.97 to 7.07 (m, 3H), 7.24 to 7.30 (m, 1H).

(d) Preparation of ethyl 3-{4-methoxy-3-[3-(3-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 1(a),
5 by reaction of 2.5 g (6.12 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f) with 1.26 g (6.73 mmol) of 3-methoxymethoxybenzyl chloride obtained above, 2.90 g (85%) of the expected
10 compound were obtained in the form of a yellow oil.
¹H NMR (CDCl₃): 1.27 to 1.35 (m, 15H), 1.69 (s, 4H), 3.43 (s, 3H), 3.77 (s, 3H), 4.24 (q, J = 7.25 Hz, 2H), 4.99 (s, 2H), 5.09 (s, 2H), 6.31 (d, J = 15.75 Hz, 1H), 6.88 to 6.96 (m, 5H), 7.16 to 7.23 (m, 2H), 7.45 to 7.51
15 (m, 2H), 7.67 (d, J = 15.75 Hz, 1H).

(e) Synthesis of 3-{4-methoxy-3-[3-(3-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h),
20 starting from 1.0 g (1.79 mmol) of ethyl 3-{4-methoxy-3-[3-(3-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 813 mg (85%) of the expected compound were obtained in the form of a white powder having a melting
25 point of 144°-146°C.

EXAMPLE 31:

Synthesis of 3-{4-methoxy-3-[3-(4-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

5 (a) Preparation of 4-methoxymethoxybenzaldehyde:

In a manner similar to that of Example 1(a), by reaction of 30 g (245 mmol) of 4-hydroxybenzaldehyde with 21 ml (270 mmol) of methoxymethyl chloride, 40.7 g
10 (100%) of the expected compound were obtained in the form of an orange oil.

¹H NMR (CDCl₃): 3.49 (s, 3H), 5.28 (s, 2H), 7.15 (d, J = 8.75 Hz, 2H), 7.84 (d, J = 8.75 Hz, 2H), 9.9 (s, 1H)

15 (b) Preparation of (4-methoxymethoxyphenyl)methanol:

In a manner similar to that of Example 29(b), by reaction of 40 g (240 mmol) of 4-methoxymethoxybenzaldehyde with 3.65 g (96 mmol) of lithium aluminum hydride, 15 g (37%) of the expected compound were
20 obtained in the form of a colorless oil.

¹H NMR (CDCl₃): 1.71 (t, J = 5.75 Hz, 1H), 3.47 (s, 3H), 4.61 (d, J = 5.75 Hz, 2H), 5.17 (s, 2H), 7.02 (d, J = 8.5 Hz, 2H), 7.29 (d, J = 8.5 Hz, 2H).

(c) Preparation of 4-methoxymethoxybenzyl chloride:

In a manner similar to that of Example 29(c), by reaction of 5.0 g (29.7 mmol) of

5 (4-methoxymethoxyphenyl)methanol with 8.27 ml (59.5 mmol) of triethylamine and 2.76 ml (35.7 mmol) of methanesulfonyl chloride, 2.66 g (48%) of the expected compound were obtained in the form of a yellow oil.
10 ^1H NMR (CDCl_3): 3.47 (s, 3H), 4.56 (s, 2H), 5.17 (s, 2H), 7.02 (d, $J = 8.75$ Hz, 2H), 7.31 (d, $J = 8.75$ Hz, 2H).

(d) Preparation of ethyl 3-{4-methoxy-3-[3-(4-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

15 In a manner similar to that of Example 1(a), by reaction of 2.5 g (6.12 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f) with 1.26 g (6.73 mmol) of 4-methoxymethoxybenzyl
20 chloride obtained above, 3.0 g (87%) of the expected compound were obtained in the form of a white powder having a melting point of $106^\circ\text{-}108^\circ\text{C}$.

(e) Synthesis of 3-{4-methoxy-3-[3-(4-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 1.0 g (1.79 mmol) of ethyl 3-{4-methoxy-3-[3-(4-methoxymethoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained
above, 877 mg (91%) of the expected compound were
obtained in the form of a white powder having a melting
10 point of 184°-186°C.

EXAMPLE 32:

Synthesis of 3-{4-methoxy-3-[3-(3-hydroxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

15 (a) Preparation of ethyl 3-{4-methoxy-3-[3-(3-hydroxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 2(g)
but in ethanol as solvent, starting from 1.9 g
20 (3.4 mmol) of ethyl 3-{4-methoxy-3-[3-(3-methoxy-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained in
Example 30(d), 1.38 g (79%) of the expected compound was

obtained in the form of a white powder having a melting point of 131°-133°C.

(b) Synthesis of 3-{4-methoxy-3-[3-(3-hydroxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 1.11 g (2.16 mmol) of ethyl 3-{4-methoxy-3-[3-(3-hydroxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 1.0 g (95%) of the expected compound was obtained in the form of a white powder having a melting point of 229°-231°C.

EXAMPLE 33:

Synthesis of 3-[4-fluoro-3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid:

In a manner similar to that of Example 1(c), by reaction of 2.0 g (8.12 mmol) of 3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid (prepared in Example 5(a) of the patent WO-97/33881) and 1.54 g (6.25 mmol) of 3-(3-bromo-4-fluorophenyl)acrylic acid, 1.45 g (63%) of the expected compound was obtained in the form of a beige powder having a melting point of 181°-183°C.

EXAMPLE 34:

Synthesis of 3-{4-hydroxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

5 (a) Preparation of 2-bromo-3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a), by reaction of 5.0 g (17.7 mmol) of 2-bromo-3-hydroxy-
10 5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene with 3.0 g (19.4 mmol) of 3-methoxybenzyl chloride, 7.14 g (100%) of the expected compound were obtained in the form of a chestnut-brown oil.

¹H NMR (CDCl₃): 1.21 (s, 6H), 1.24 (s, 6H), 1.64 (s, 4H), 3.82 (s, 3H), 5.09 (s, 2H), 6.83 to 6.87 (m, 2H),
15 7.03 to 7.08 (m, 2H), 7.29 (m, 1H), 7.43 (s, 1H).

(b) Preparation of 3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid:

20 In a manner similar to that of Example 1(b), starting from 7.10 g (17.6 mmol) of 2-bromo-3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene obtained above, 4.85 g (74%) of

the expected compound were obtained in the form of a slightly tacky beige powder.

¹H NMR (CDCl₃): 1.26 (s, 6H), 1.29 (s, 6H), 1.67 (s, 4H), 3.81 (s, 3H), 5.06 (s, 2H), 6.85 to 7.0 (m, 3H),
5 7.31 (t, J = 7.75 Hz, 1H), 7.78 (s, 1H).

(c) Preparation of ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate:

In a manner similar to that of Example 2(c), by reaction of 30 g (122 mmol) of ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate obtained above with
10 30.27 g (135 mmol) of triethyl phosphonoacetate, 34.6 g (90%) of the expected compound were obtained in the form of white crystals having a melting point of 64°-66°C.

(d) Preparation of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate:

In a manner similar to that of Example 2(g), starting from 30 g (95 mmol) of ethyl 3-(3-bromo-4-methoxymethoxyphenyl)acrylate obtained above, 25.8 g (100%) of the expected compound were obtained in the
20 form of a white powder having a melting point of 113°-115°C.

(e) Preparation of ethyl 3-{4-hydroxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 1(c),
5 by reaction of 1.0 g (3.7 mmol) of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate obtained above with 1.77 g
(4.8 mmol) of 3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid
obtained in Example 34(c), 630 mg (33%) of the expected
10 compound were obtained in the form of a colorless oil.
¹H NMR (CDCl₃): 1.25 to 1.36 (m, 15H), 1.70 (s, 4H),
3.78 (s, 3H), 4.26 (q, J = 7.25 Hz, 2H), 4.99 (s, 2H),
6.32 (d, J = 15.75 Hz, 1H), 6.84 (d, J = 8.5 Hz, 2H),
6.93 (s, 1H), 6.98 to 7.02 (m, 2H), 7.19 (d, J = 8.5 Hz,
15 1H), 7.25 (s, 1H), 7.40 (d, J = 2.25 Hz, 1H), 7.69 (d, J
= 15.75 Hz, 1H).

(f) Synthesis of 3-{4-hydroxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}-acrylic acid:

20 In a manner similar to that of Example 2(h),
starting from 600 mg (1.16 mmol) of ethyl 3-{4-hydroxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above,
539 mg (95%) of the expected compound were obtained in
25 the form of a white powder having a melting point of
197°-199°C.

EXAMPLE 35:

Synthesis of 3-{4-hydroxy-3-[3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

5 (a) Preparation of 2-bromo-3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a), by reaction of 5.0 g (17.7 mmol) of 3-hydroxy-2-bromo-10 5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene with 3.0 g (19.4 mmol) of 4-methoxybenzyl chloride, 7.14 g (100%) of the expected compound were obtained in the form of a chestnut-brown oil.

¹H NMR (CDCl₃): 1.21 (s, 6h), 1.24 (s, 6H), 1.64 (s, 15 4H), 3.81 (s, 3H), 5.04 (s, 2H), 6.83 (s, 1H), 6.92 (d, J = 7.5 Hz, 2H), 7.39 (d, J = 7.5 Hz, 2H), 7.43 (s, 1H).

(b) Preparation of 3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid:

20 In a manner similar to that of Example 1(b), starting from 7.14 g (17.7 mmol) of 2-bromo-3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene obtained above, 3.55 g (54%) of

the expected compound were obtained in the form of an orange oil.

¹H (CDCl₃): 1.28 (s, 12H), 1.57 (s, 4H), 3.82 (s, 3H),
5.03 (s, 2H), 5.78 (s, 2H), 6.89 (s, 1H), 6.93 (d,
5 J = 8.75 Hz, 1H), 7.35 (d, J = 8.5 Hz, 1H), 7.78 (s,
1H).

(c) Preparation of ethyl 3-{4-hydroxy-3-[3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

10 In a manner similar to that of Example 1(c),
by reaction of 1.0 g (3.7 mmol) of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate obtained in Example 34(d) with
2.0 g (5.5 mmol) of 3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid
15 obtained above, 930 mg (49%) of the expected compound
were obtained in the form of a colorless oil.

¹H NMR (CDCl₃): 1.29 (s, 12H), 1.34 (t, J = 7.0 Hz, 3H),
1.70 (s, 4H), 3.72 (s, 3H), 4.26 (q, J = 7.25 Hz, 2H),
5.02 (s, 2H), 6.33 (d, J = 15.75 Hz, 1H), 6.70 (s, 1H),
20 6.80 to 6.87 (m, 3H), 7.02 (d, J = 7.25 Hz, 2H), 7.18 to
7.25 (m, 2H), 7.42 (d, J = 2.25 Hz, 1H), 7.48 (d,
J = 8.5 Hz, 1H), 7.69 (d, J = 16.0 Hz, 1H).

(d) Synthesis of 3-{4-hydroxy-3-[3-(3-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 930 mg (1.8 mmol) of ethyl 3-{4-hydroxy-3-[3-(4-methoxybenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above,
800 mg (91%) of the expected compound were obtained in
the form of a white powder having a melting point of
10 170°-172°C.

EXAMPLE 36:

Synthesis of 3-{4-hydroxy-3-[3-(4-fluorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

15 (a) Preparation of 2-bromo-3-(4-fluorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a),
by reaction of 5.0 g (17.7 mmol) of 3-hydroxy-2-bromo-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene with
20 3.67 g (19.4 mmol) of 4-fluorobenzyl bromide, 4.3 g
(62%) of the expected compound were obtained in the form
of light chestnut-brown crystals having a melting point
of 74°-76°C.

(b) Preparation of 3-(4-fluorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid:

In a manner similar to that of Example 1(b),
5 starting from 4.3 g (11.0 mmol) of 2-bromo-3-(4-fluorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene obtained above, 3.24 g (83%) of the expected compound were obtained in the form of an orange oil.

10 ¹H NMR (CDCl₃): 1.27 (s, 6H), 1.29 (s, 6H), 1.67 (s, 4H), 5.08 (s, 2H), 5.81 (s, 2H), 7.06 to 7.13 (m, 2H), 7.38 to 7.43 (m, 2H), 7.80 (s, 1H).

(c) Preparation of ethyl 3-{4-hydroxy-3-[3-(4-fluorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar of Example 1(c), by
reaction of 1.0 g (3.7 mmol) of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate obtained in Example 34(d) with
1.97 g (5.5 mmol) of 3-(4-fluorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid
20 obtained above, 825 mg (44%) of the expected compound were obtained in the form of a white solid having a melting point of 123°-125°C.

(d) Synthesis of 3-{4-hydroxy-3-[3-(3-fluorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 800 mg (1.6 mmol) of ethyl 3-{4-hydroxy-3-[3-(4-fluorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 425 mg (56%) of the expected compound were obtained in the form of a white powder having a melting point of
10 157°-159°C.

EXAMPLE 37:

Synthesis of 3-{4-hydroxy-3-[3-(4-chlorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

15 (a) Preparation of 2-bromo-3-(4-chlorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a),
by reaction of 5.0 g (17.7 mmol) of 3-hydroxy-2-bromo-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene with
20 3.12 g (19.4 mmol) of 4-chlorobenzyl chloride, 7.22 g (100%) of the expected compound were obtained in the form of pale chestnut-brown crystals having a melting point of 108°-110°C.

(b) Preparation of 3-(4-chlorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid:

In a manner similar to that of Example 1(b),
5 starting from 7.14 g (17.5 mmol) of 2-bromo-3-(4-chlorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene obtained above, 3.50 g (54%) of the expected compound were obtained in the form of an orange oil.

10 ¹H NMR (CDCl₃): 1.26 (s, 6H), 1.29 (s, 6H), 1.67 (s, 4H), 5.09 (s, 2H), 5.72 (s, 2H), 6.84 (s, 1H), 7.37 (m, 4H), 7.80 (s, 1H).

(c) Preparation of ethyl 3-{4-hydroxy-3-[3-(4-chlorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 1(c),
by reaction of 1.0 g (3.7 mmol) of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate obtained in Example 34(d) with
2.05 g (5.5 mmol) of 3-(4-chlorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid
20 obtained above, 590 mg (30%) of the expected compound were obtained in the form of a white solid having a melting point of 152°-154°C.

(d) Synthesis of 3-{4-hydroxy-3-[3-(4-chlorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 580 mg (1.12 mmol) of ethyl 3-{4-hydroxy-3-[3-(4-chlorobenzyloxy)-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above,
382 mg (70%) of the expected compound were obtained in
the form of a white powder having a melting point of
10 147°-149°C.

EXAMPLE 38:

Synthesis of 3-[4-hydroxy-3-(3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid:

15 (a) Preparation of ethyl 3-(4-hydroxy-3-[3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 1(c),
by reaction of 1.0 g (3.7 mmol) of ethyl 3-(3-bromo-4-
20 hydroxyphenyl)acrylate obtained in Example 34(d) with
1.6 g (5.5 mmol) of 3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid
obtained in Example 13(b), 150 mg (9%) of the expected

compound were obtained in the form of a white solid having a melting point of 101°-103°C.

(b) Synthesis of 3-{4-hydroxy-3-[3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h), starting from 150 mg (0.34 mmol) of ethyl 3-{4-hydroxy-3-[3-methoxymethoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 100 mg (72%) of the expected compound were obtained in the form of a white powder having a melting point of 191°-193°C.

EXAMPLE 39:

Synthesis of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-propyloxy-3-biphenyl)acrylic acid:

(a) Preparation of 1-bromo-3,5-di-tert-butyl-2-propyloxybenzene:

In a manner similar to that of Example 1(a), by reaction of 15 g (52.6 mmol) of 6-bromo-2,4-di-tert-butylphenol with 7.4 g (57.9 mmol) of 1-bromopropane, 17.2 g (100%) of the expected compound were obtained in the form of orange crystals having a melting point of 41°-43°C.

(b) Preparation of 2-propyloxy-3,5-di-tert-butylphenylboronic acid:

In a manner similar to that of Example 1(b), starting from 16.0 g (48.9 mmol) of 1-bromo-3,5-di-tert-butyl-2-propyloxybenzene obtained above, 9.1 g (63%) of the expected compound were obtained in the form of a white powder having a melting point of 104°-106°C.

(c) Preparation of ethyl 3-(3',5'-di-tert-butyl-6-hydroxy-2'-propyloxy-3-biphenyl)acrylate:

In a manner similar to that of Example 1(c), by reaction of 1.0 g (3.7 mmol) of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate obtained in Example 34(d) with 2.14 g (7.33 mmol) of 2-propyloxy-3,5-di-tert-butylphenylboronic acid obtained above, 70 mg (4%) of the expected compound were obtained in the form of a colorless oil.

¹H NMR (CDCl₃): 0.85 (t, J = 7.5 Hz, 3H), 1.34 (s, 9H), 1.40 (t, J = 7.75 Hz, 3H), 1.45 (s, 9H), 1.55 to 1.59 (m, 2H), 3.43 to 3.65 (m, 2H), 4.27 (q, J = 8.25 Hz, 2H), 6.37 (d, J = 15.75 Hz, 1H), 7.05 (d, J = 8.25 Hz, 1H), 7.16 (d, J = 2.5 Hz, 1H), 7.30 (s, 1H), 7.43 (d, J = 2.5 Hz, 1H), 7.50 to 7.55 (m, 2H), 7.72 (d, J = 16.0 Hz, 1H).

(d) Synthesis of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-propyloxy-3-biphenyl)acrylic acid:

In a manner similar to that of Example 2(h), starting from 70 mg (0.16 mmol) of ethyl 3-(3',5'-di-
5 tert-butyl-6-hydroxy-2'-propyloxy-3-biphenyl)acrylate obtained above, 50 mg (75%) of the expected compound were obtained in the form of a white powder having a melting point of 187°-189°C.

EXAMPLE 40:

10 Synthesis of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-butyloxy-3-biphenyl)acrylic acid:

(a) Preparation of 1-bromo-3,5-di-tert-butyl-2-butyloxybenzene:

In a manner similar to that of Example 1(a),
15 by reaction of 15 g (52.6 mmol) of 2-bromo-4,6-di-tert-butylphenol with 7.93 g (57.9 mmol) of 1-bromobutane, 17.7 g (100%) of the expected compound were obtained in the form of an orange oil.

¹H NMR (CDCl₃): 0.99 (t, J = 7.5 Hz, 3H), 1.28 (s, 9H),
20 1.39 (s, 9H), 1.45 to 1.60 (m, 2H), 1.80 to 1.92 (m, 2H), 4.03 (t, J = 7.0 Hz, 2H), 7.27 (d, J = 2.5 Hz, 1H), 7.39 (d, J = 2.5 Hz, 1H).

(b) Preparation of 2-butyloxy-3,5-di-tert-butylphenylboronic acid:

In a manner similar to that of Example 1(b), starting from 16.0 g (46.8 mmol) of 1-bromo-3,5-di-tert-butyl-2-butyloxybenzene obtained above, 14 g (97%) of the expected compound were obtained in the form of a white powder having a melting point of 92°-94°C.

(c) Preparation of ethyl 3-(3',5'-di-tert-butyl-6-hydroxy-2'-butyloxy-3-biphenyl)acrylate:

In a manner similar to that of Example 1(c), by reaction of 6.2 g (22.8 mmol) of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate obtained in Example 34(d) with 14 g (45.7 mmol) of 2-butyloxy-3,5-di-tert-butylphenylboronic acid obtained above, 700 mg (7%) of the expected compound were obtained in the form of a yellow oil.

¹H NMR (CDCl₃): 0.78 (t, J = 7.5 Hz, 3H), 1.31 to 1.56 (m, 25H), 3.20 to 3.66 (m, 2H), 4.27 (q, J = 7.25 Hz, 2H), 6.37 (d, J = 16.0 Hz, 1H), 7.06 (d, J = 8.25 Hz, 1H), 7.16 (d, J = 2.5 Hz, 1H), 7.30 (s, 1H), 7.43 (d, J = 2.25 Hz, 1H), 7.49 to 7.55 (m, 2H), 7.72 (d, J = 16.0 Hz, 1H).

(d) Synthesis of 3-(3',5'-di-tert-butyl-6-hydroxy-2'-butyloxy-3-biphenyl)acrylic acid:

In a manner similar to that of Example 2(h), starting from 430 mg (0.95 mmol) of ethyl 3-(3',5'-di-tert-butyl-6-hydroxy-2'-butyloxy-3-biphenyl)acrylate
5 obtained above, 300 mg (74%) of the expected compound were obtained in the form of a white powder having a melting point of 194°-196°C.

EXAMPLE 41:

10 Synthesis of 3-(2'-butoxy-3',5'-di-tert-butyl-6-methoxy-3-biphenyl)acrylic acid:

(a) Preparation of ethyl 3-(2'-butoxy-3',5'-di-tert-butyl-6-methoxy-3-biphenyl)acrylate:

In a manner similar to that of Example 1(a),
15 by reaction of 230 mg (0.5 mmol) of ethyl 3-(3',5'-di-tert-butyl-6-hydroxy-2'-butyloxy-3-biphenyl)acrylate obtained in Example 40(c) with 82 mg (0.58 mmol) of iodomethane, 230 mg (100%) of the expected compound were obtained in the form of a yellow oil.

20 ¹H NMR (CDCl₃): 0.67 (t, J = 7.25 Hz, 3H), 1.07 (m, 2H), 1.25 to 1.36 (m, 14H), 1.43 (s, 9H), 3.40 (br, 2H), 3.84 (s, 3H), 4.25 (q, J = 7.0 Hz, 2H), 6.33 (d, J = 15.5 Hz, 1H), 6.98 (d, J = 8.5 Hz, 1H), 7.11 (d, J = 2.25 Hz, 1H), 7.35 (d, J = 2.5 Hz, 1H), 7.48 (dd, J₁ = 8.5 Hz,

J2 = 2.25 Hz, 1H), 7.56 (d, J = 2.25 Hz, 1H), 7.68 (d, J = 15.75 Hz, 1H).

(b) Synthesis of 3-(2'-butoxy-3',5'-di-tert-butyl-6-methoxy-3-biphenyl)acrylic acid:

5 In a manner similar to that of Example 2(h), starting from 230 mg (0.51 mmol) of ethyl 3-(2'-butoxy-3',5'-di-tert-butyl-6-methoxy-3-biphenyl)acrylate obtained above, 120 mg (55%) of the expected compound were obtained in the form of a white powder having a
10 melting point of 204°-206°C.

EXAMPLE 42:

Synthesis of 3-(3',5'-di-tert-butyl-6-methoxy-2'-propoxy-3-biphenyl)acrylic acid:

(a) Preparation of ethyl 3-(3',5'-di-tert-butyl-6-methoxy-2'-propoxy-3-biphenyl)acrylate:
15

 In a manner similar to that of Example 1(c), by reaction of 1.0 g (3.5 mmol) of ethyl 3-(3-bromo-4-methoxyphenyl)acrylate obtained in Example 13(d) with 1.54 g (5.26 mmol) of 2-propyloxy-3,5-di-tert-butylphenylboronic acid obtained in Example 39(b),
20 1.20 g (76%) of the expected compound was obtained in the form of a white powder having a melting point of 107°-109°C.

(b) Synthesis of 3-(3',5'-di-tert-butyl-6-methoxy-2'-propoxy-3-biphenyl)acrylic acid:

In a manner similar to that of Example 2(h), starting from 1.0 g (2.2 mmol) of ethyl 3-(3',5'-di-
5 tert-butyl-6-methoxy-2'-propoxy-3-biphenyl)acrylate obtained above, 800 mg (85%) of the expected compound were obtained in the form of a white powder having a melting point of 207°-209°C.

EXAMPLE 43:

10 Synthesis of 3-[4-hydroxy-3-(5,5,8,8-tetramethyl-4-propoxy-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid:

(a) Preparation of 3-bromo-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphth-1-ol:

15 9.73 g (73 mmol) of aluminum chloride were introduced together with 80 ml of dichloromethane under a stream of nitrogen into a 2 liter three-necked flask, and a solution of 25 g (145 mmol) of 3-bromophenol in 100 ml of dichloromethane was added dropwise, and the
20 mixture was stirred at room temperature for 15 minutes. Then a solution of 26.2 g (143 mmol) of 2,5-dichloro-2,5-dimethylhexane in 100 ml of dichloromethane was introduced and the mixture was stirred at room temperature for 1 hour. Then a solution of 40 g

(218 mmol) of 2,5-dichloro-2,5-dimethylhexane in 200 ml of dichloromethane was added in four portions with one hour of stirring between each addition and the mixture was stirred for 15 hours. The dichloromethane was
5 evaporated from the reaction medium, the residue was taken up in ethyl acetate and water, and the organic phase was decanted, dried over magnesium sulfate, filtered and evaporated. The residue obtained was purified by chromatography on silica, eluting first with
10 heptane and then with a 95:5 heptane/ethyl acetate mixture. After evaporation of the solvents, 30 g (73%) of the expected compound were obtained in the form of oily black crystals.
¹H NMR (CDCl₃): 1.25 (s, 6H), 1.38 (s, 6H), 1.58 to 1.67
15 (m, 4H), 4.99 (s, 1H), 6.25 (d, J = 2.0 Hz, 1H), 7.04 (d, J = 2.0 Hz, 1H).

(b) Preparation of 2-bromo-4-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene:

In a manner similar to that of Example 1(a),
20 by reaction of 18 g (63.6 mmol) of 3-bromo-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphth-1-ol obtained in the preceding example with 8.61 g (70 mmol) of 1-bromopropane, 16.2 g (78%) of the expected compound were obtained in the form of yellow crystals having a melting
25 point of 86°-88°C.

(c) Preparation of 4-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid:

In a manner similar to that of Example 1(b), starting from 16.0 g (49.2 mmol) of 2-bromo-4-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalene
5 obtained above, 6.2 g (46%) of the expected compound were obtained in the form of a white powder having a melting point of 230°-232°C.

(d) Preparation of ethyl 3-[4-hydroxy-3-(5,5,8,8-tetramethyl-4-propoxy-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylate:

In a manner similar to that of Example 1(c), by reaction of 1.0 g (3.7 mmol) of ethyl 3-(3-bromo-4-hydroxyphenyl)acrylate obtained in Example 34(d) with
15 1.61 g (5.5 mmol) of 4-propyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid obtained in Example 43(c), 640 mg (40%) of the expected compound were obtained in the form of a white powder having a melting point of 114°-116°C.

(e) Synthesis of 3-[4-hydroxy-3-(5,5,8,8-tetramethyl-4-propoxy-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylic acid:

In a manner similar to that of Example 2(h), starting from 640 mg (1.47 mmol) of ethyl 3-[4-hydroxy-

3-(5,5,8,8-tetramethyl-4-propoxy-5,6,7,8-tetrahydro-2-naphthyl)phenyl]acrylate obtained above, 270 mg (45%) of the expected compound were obtained in the form of a white powder having a melting point of 209°-211°C.

5

EXAMPLE 44:

Synthesis of 3-{4-methoxy-3-[3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

10 (a) Preparation of ethyl 3-{4-methoxy-3-[3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate:

In a manner similar to that of Example 1(a), by reaction of 1.40 g (3.4 mmol) of ethyl 3-[3-(3-hydroxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl)-4-methoxyphenyl]acrylate obtained in Example 13(f) with 450 μ l (3.77 mmol) of benzyl bromide, 1.64 g (97%) of the expected compound was obtained in the form of a yellow oil.

¹H NMR (CDCl₃): 1.28 to 1.35 (m, 15H), 1.69 (s, 4H), 3.76 (s, 3H), 4.25 (q, J = 7.0 Hz, 2H), 5.02 (s, 2H), 6.30 (d, J = 15.75 Hz, 1H), 6.89 (s, 1H), 6.93 (d, J = 8.25 Hz, 1H), 7.19 (s, 1H), 7.20 to 7.35 (m, 5H), 7.43 to 7.52 (m, 2H), 7.67 (d, J = 16.0 Hz, 1H).

(b) Synthesis of 3-{4-methoxy-3-[3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylic acid:

In a manner similar to that of Example 2(h),
5 starting from 1.64 g (3.3 mmol) of ethyl 3-{4-methoxy-3-[3-benzyloxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-2-naphthyl]phenyl}acrylate obtained above, 1.27 g (82%) of the expected compound were obtained in the form of a white powder having a melting point of 200°-202°C.

10

EXAMPLE 45:

Synthesis of 3-methyl-5-[3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]penta-2,4-dienoic acid:

(a) Preparation of 3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)benzaldehyde:
15

In a manner similar to that of Example 1(c),
by reaction of 2.0 g (8.12 mmol) of 3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthylboronic acid (prepared in Example 5(a) of the patent WO 97/33881) and
20 1.89 g (10.15 mmol) of 3-bromobenzaldehyde, 3.10 g (100%) of the expected compound were obtained in the form of a white powder having a melting point of 99°-101°C.

(b) Preparation of ethyl 3-methyl-5-[3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]penta-2,4-dienoate:

In a manner similar to that of Example 2(c),
5 by reaction of 2.0 g (6.53 mmol) of 3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)benzaldehyde obtained in the preceding example and 1.90 g (7.20 mmol) of triethyl 3-methyl-4-phosphonocrotonate, 1.80 g (66%) of the expected compound was obtained in the form of a
10 white powder having a melting point of 83°-85°C.

(c) Synthesis of 3-methyl-5-[3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]penta-2,4-dienoic acid:

In a manner similar to that of Example 2(h),
15 starting from 1.00 g (2.4 mmol) of ethyl 3-methyl-5-[3-(3,5,5,8,8-pentamethyl-5,6,7,8-tetrahydro-2-naphthyl)phenyl]penta-2,4-dienoate obtained above and after separation of the isomers on a fine silica column (7-35 μ m) eluted with a 95:5 heptane/ethyl acetate
20 mixture, 420 mg (45%) of the expected compound were obtained in the form of a white powder having a melting point of 218°-220°C.

B. BIOLOGICAL TEST EXAMPLES:

The biological tests carried out correspond to those described above. The results obtained are reported in the following Table:

TABLE:

Compound No.	RXR binding	RXR α transactivation	
	K _O (nM)	Agonist AC ₅₀ (nM)	Antagonist IC ₅₀ (nM)
1	394	384	-----
2	23	inactive	920
3	10	inactive	729
4	17	165	-----
5	76	inactive	4000
6	14	inactive	505
7	153	32	-----
8	235	inactive	1100
9	231	inactive	631

C. EXAMPLES OF SPECIFIC FORMULATIONS BASED ON THE COMPOUNDS OF THE INVENTION:

(1) ORAL ROUTE:

(a) The following composition was formulated as a 0.8 g tablet:

	Compound of Example 3	0.005 g
	Pregelatinized starch	0.265 g
	Microcrystalline cellulose	0.300 g
	Lactose	0.200 g
5	Magnesium stearate	0.030 g

For the treatment of acne, 1 to 3 tablets are administered to an adult individual per day for 3 to 6 months, depending on the severity of the condition treated.

10 (b) A drinkable suspension for packaging in 5 ml vials was formulated:

	Compound of Example 2	0.050 g
	Glycerol	0.500 g
	70% sorbitol	0.500 g
15	Sodium saccharinate	0.010 g
	Methyl parahydroxybenzoate	0.040 g
	Flavoring	qs
	Purified water	qs 5 ml

20 For the treatment of acne, 1 vial is administered to an adult individual per day for 3 months, depending on the severity of the condition treated.

(c) The following composition for packaging in gelatin capsules was formulated:

Compound of Example 5	0.025 g
Corn starch	0.060 g
Lactose qs	0.300 g

5 The gelatin capsules comprised gelatin,
titanium dioxide and a preservative.

For the treatment of psoriasis, 1 gelatin capsule is administered to an adult individual per day for 30 days.

(2) TOPICAL ROUTE:

10 (a) The following nonionic water-in-oil cream
was formulated:

Compound of Example 6	0.100 g
Mixture of emulsifying lanolin alcohols, waxes and refined oils, 15 marketed by BDF under the trademark "anhydrous Eucerin"	39.900 g
Methyl para-hydroxybenzoate	0.075 g
Propyl para-hydroxybenzoate	0.075 g
Sterile demineralized water qs	100.000 g

20 This cream is applied to psoriatic skin once
or twice a day for 30 days.

(b) A gel was formulated from the following composition:

Compound of Example 18	0.050 g
25 Base erythromycin	4.000 g

Butylhydroxytoluene 0.050 g
Hydroxypropylcellulose marketed
by Hercules under the trademark
"Klucel HF" 2.000 g
5. Ethanol (at 95°) qs 100.000 g

This gel is applied to skin afflicted with
dermatitis or acneic skin 1 to 3 times a day for 6 to
12 weeks, depending on the severity of the condition
treated.

10 (c) An anti-seborrhoeic lotion was formulated
by mixing together the following ingredients:

Compound of Example 4 0.030 g
Propylene glycol 5.000 g
Butylhydroxytoluene 0.100 g
15 Ethanol (at 95°) qs 100.000 g

This lotion is applied twice a day to a
seborrhoeic scalp, and a significant improvement is
observed within a period of between 2 and 6 weeks.

20 (d) A cosmetic composition to combat the
harmful effects of the sun was formulated by mixing
together the following ingredients:

Compound of Example 25 1.000 g
Benzylidenecamphor 4.000 g
Fatty acid triglycerides 31.000 g
25 Glyceryl monostearate 6.000 g
Stearic acid 2.000 g

	Cetyl alcohol	1.200 g
	Lanolin	4.000 g
	Preservatives	0.300 g
	Propylene glycol	2.000 g
5	Triethanolamine	0.500 g
	Fragrance	0.400 g
	Demineralized water qs	100.000 g

This composition is applied daily to combat photoinduced skin aging.

10 (e) The following nonionic oil-in-water cream was formulated:

	Compound of Example 38	0.500 g
	Vitamin D3	0.020 g
	Cetyl alcohol	4.000 g
15	Glyceryl monostearate	2.500 g
	PEG-50 stearate	2.500 g
	Karite butter	9.200 g
	Propylene glycol	2.000 g
	Methyl para-hydroxybenzoate	0.075 g
20	Propyl para-hydroxybenzoate	0.075 g
	Sterile demineralized water qs	100.000 g

This cream is applied to psoriatic skin once or twice a day for 30 days.

(f) A topical gel was formulated by mixing
25 together the following ingredients:

	Compound of Example 19	0.050 g
	Ethanol	43.000 g
	α -tocopherol	0.050 g
	Carboxyvinyl polymer marketed	
5	under the trademark "Carbopol 941"	
	by Goodrich	0.500 g
	Triethanolamine as an aqueous	
	solution at 20% by weight	3.800 g
	Water	9.300 g
10	Propylene glycol qs	100.000 g

This gel is applied in the treatment of acne 1
to 3 times a day for 6 to 12 weeks, depending on the
severity of the condition treated.

(g) A hair lotion to combat hair loss and to
15 promote regrowth of the hair was formulated by mixing
together the following ingredients:

	Compound of Example 41	0.05 g
	Compound marketed under the	
	trademark "Minoxidil"	1.00 g
20	Propylene glycol	20.00 g
	Ethanol	34.92 g
	Polyethylene glycol (molecular	
	weight = 400)	40.00 g
	Butylhydroxyanisole	0.01 g
25	Butylhydroxytoluene	0.02 g
	Water qs	100.00 g

This lotion is applied twice a day for 3 months to a scalp which has suffered considerable hair loss.

(h) An anti-acne cream was formulated by mixing together the following ingredients:

	Compound of Example 13	0.050 g
	Retinoic acid	0.010 g
	Mixture of glycerol stearate and polyethylene glycol stearate (75 mol),	
10	marketed under the trademark "Gelot 64" by Gattefosse	15.000 g
	Kernel oil polyoxyethylenated with 6 mol of ethylene oxide, marketed under the trademark	
15	"Labrafil M2130 CS" by Gattefosse	8.000 g
	Perhydrosqualene	10.000 g
	Preservatives	qs
	Polyethylene glycol (molecular weight = 400)	8.000 g
20	Disodium salt of ethylenediamine-tetraacetic acid	0.050 g
	Purified water qs	100.000 g

This cream is applied to skin afflicted with dermatitis or to acneic skin 1 to 3 times a day for 6 to 12 weeks.

(i) An oil-in-water cream was formulated from the following composition:

	Compound of Example 29	0.020 g
	Betamethasone 17-valerate	0.050 g
5	S-Carboxymethylcysteine	3.000 g
	Polyoxyethylene stearate (40 mol of ethylene oxide) marketed under the trademark "Myrj 52" by Atlas	4.000 g
10	Sorbitan monolaurate, polyoxyethylene with 20 mol of ethylene oxide, marketed under the trademark "Tween 20" by Atlas	1.800 g
	Mixture of glyceryl mono- and distearate marketed under the trademark "Geleol" by Gattefosse . .	4.200 g
15	Propylene glycol	10.000 g
	Butylhydroxyanisole	0.010 g
	Butylhydroxytoluene	0.020 g
	Cetostearyl alcohol	6.200 g
20	Preservatives	qs
	Perhydrosqualene	18.000 g
	Mixture of caprylic/capric triglycerides marketed under the trademark "Miglyol 812" by Dynamit Nobel	4.000 g
25	Triethanolamine (99% by weight) . .	2.500 g
	Water qs	100.000 g

This cream is applied twice a day to skin afflicted with dermatitis, for 30 days.

5	(j) The following oil-in-water cream was formulated:		
	Lactic acid	5.000 g	
	Compound of Example 33	0.020 g	
	Polyoxyethylene stearate (40 mol of ethylene oxide) marketed under the trademark "Myrj 52" by Atlas	4.000 g	
	Sorbitan monolaurate, polyoxyethylene with 20 mol of ethylene oxide, marketed under the trademark "Tween 20" by Atlas	1.800 g	
	Mixture of glyceryl mono- and distearate marketed under the trademark "Geleol" by Gattefosse.	4.200 g	
	Propylene glycol	10.000 g	
	Butylhydroxyanisole	0.010 g	
	Butylhydroxytoluene	0.020 g	
	Cetostearyl alcohol	6.200 g	
	Preservatives	qs	
	Perhydrosqualene	18.000 g	
	Mixture of caprylic/capric triglycerides marketed under the trademark "Miglyol 812" by Dynamit Nobel	4.000 g	
	Water qs	100.000 g	

This cream is applied once a day to combat aging, whether photoinduced or chronological aging.

While the invention has been described in terms of various preferred embodiments, the skilled
5 artisan will appreciate that various modifications, substitutions, omissions, and changes may be made without departing from the spirit thereof. Accordingly, it is intended that the scope of the present invention be limited solely by the scope of the following claims,
10 including equivalents thereof.